

## FUSED-RING COMPOUNDS AND USE THEREOF AS DRUGS

This application is a divisional of copending U.S. Patent Application No. 09/939,374, filed August 24, 2001, which is a  
5 continuation-in-part of PCT/JP00/09181 filed on December 22, 2000.

### Technical Field

The present invention relates to a novel fused ring  
10 compound and a pharmaceutically acceptable salt thereof useful as a therapeutic agent for hepatitis C, and to an intermediate compound for the synthesis thereof. The present invention also relates to a novel use of a certain fused ring compound or a pharmaceutically acceptable salt thereof as a therapeutic agent  
15 for hepatitis C. More particularly, the present invention relates to a therapeutic agent for hepatitis C, which contains a novel fused ring compound or a Pharmaceutically acceptable salt thereof, which is effective for the prophylaxis or treatment of hepatitis C and which shows anti-hepatitis C virus (HCV)  
20 activity, particularly anti-HCV activity based on an RNA-dependent RNA polymerase inhibitory activity.

### Background Art

In 1989, a main causative virus of non-A non-B posttrans-  
25 fusion hepatitis was found and named hepatitis C virus (HCV). Since then, several types of hepatitis viruses have been found besides type A, type B and type C, wherein hepatitis caused by HCV is called hepatitis C.

The patients infected with HCV are considered to involve  
30 several percent of the world population, and the infection with HCV characteristically becomes chronic.

HCV is an envelope RNA virus, wherein the genome is a single strand plus-strand RNA, and belongs to the genus Hepacivirus of Flavivirus (from The International Committee on  
35 Taxonomy of Viruses, International Union of Microbiological Societies). Of the same hepatitis viruses, for example, hepatitis B virus (HBV), which is a DNA virus, is eliminated by the immune system and the infection with this virus ends in an acute infection except for neonates and infants having yet immature  
40 immunological competence. In contrast, HCV somehow avoids the immune system of the host due to an unknown mechanism. Once

infected with this virus, even an adult having a mature immune system frequently develops persistent infection.

When chronic hepatitis is associated with the persistent infection with HCV, it advances to cirrhosis or hepatic cancer in a high rate. ENUCLEATION of tumor by operation does not help much, because the patient often develops recurrent hepatic cancer due to the sequela inflammation in non-cancerous parts.

Thus, an effective therapeutic method of hepatitis C is desired. Apart from the symptomatic therapy to suppress inflammation with an anti-inflammatory agent, the development of a therapeutic agent that reduces HCV to a low level free from inflammation and that eradicates HCV has been strongly demanded.

At present, a treatment with interferon is the only effective method known for the eradication of HCV. However, interferon can eradicate the virus only in about one-third of the patient population. For the rest of the patients, it has no effect or provides only a temporary effect. Therefore, an anti-HCV drug to be used in the place of or concurrently with interferon is awaited in great expectation.

In recent years, Ribavirin (1- $\beta$ -D-ribofuranosyl-1H-1,2,4-triazole-3-carboxamide) has become commercially available as a therapeutic agent for hepatitis C, which is to be used concurrently with interferon. It enhances the efficacy of interferon but only to a low efficacy rate, and a different novel therapeutic agent for hepatitis C is desired.

Also, an attempt has been made to potentiate the immunocompetence of the patient with an interferon agonist, an interleukin-12 agonist and the like, thereby to eradicate the virus, but an effective pharmaceutical agent has not been found yet.

In addition, the inhibition of HCV growth, wherein HCV-specific protein is targeted, has been drawing attention these days.

The gene of HCV encodes a protein such as serine protease, RNA helicase, RNA-dependent RNA polymerase and the like. These proteins function as a specific protein essential for the growth of HCV.

One of the specific proteins, RNA-dependent RNA polymerase (hereinafter to be also briefly referred to as an HCV polymerase), is an enzyme essential for the growth of the virus. The gene replication of HCV having a plus-strand RNA gene is considered to involve synthesis of a complementary minus-strand RNA by the use of the plus-strand RNA as a template, and, using the obtained minus-strand RNA as a template, amplifying the plus-strand RNA. The portion called NS5B of a protein precursor, that HCV codes for, has been found to show an RNA-dependent RNA polymerase activity (EMBO J., 15, 12-22, 1996), and is considered to play a central role in the HCV gene replication.

Therefore, an HCV polymerase inhibitor can be a target in the development of an anti-HCV drug, and the development thereof is eagerly awaited. However, an effective HCV polymerase inhibitor has not been developed yet, like in other attempts to develop an anti-HCV drug based on other action mechanisms. As the situation stands, no pharmaceutical agent can treat hepatitis C satisfactorily.

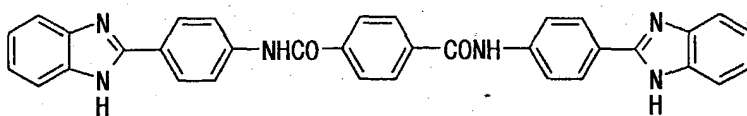
The following discloses known compounds relatively similar to the compound of the present invention.

A known therapeutic agent for hepatitis C having a benzimidazole skeleton is disclosed in WO97/36866, Japanese Patent Application under PCT laid-open under kohyo No. 2000-511899 (EP906097) and WO99/51619.

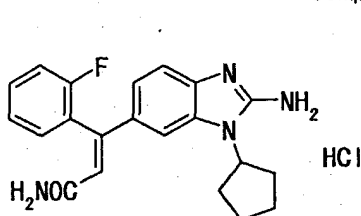
WO97/36866 discloses the following compound D and the like, and HCV helicase inhibitory activity of the compounds.

Japanese Patent Application under PCT laid-open under kohyo No. 2000-511899 (EP906097) discloses the following compound E and the like, and WO99/51619 discloses the following compound F and the like, in both of which a possibility of these compounds being effective as an HCV inhibitor is mentioned.

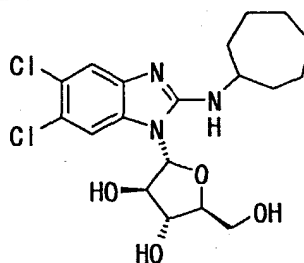
However, these publications do not include the compound disclosed in the present specification, or a disclosure suggestive thereof.



compound D



compound E

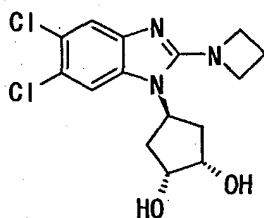


compound F

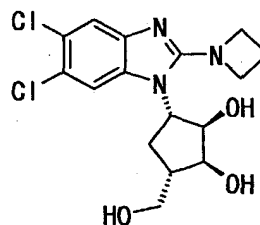
A known anti-hepatitis virus agent having a benzimidazole skeleton is disclosed in Japanese Patent Application under PCT laid-open under kohyo No. 2000-503017 (WO97/25316) and Japanese Patent Application under PCT laid-open under kohyo No. 10-505092 (WO96/7646).

WO97/25316 discloses the following compound A and the like, wherein the use thereof is for a treatment of viral infection. The target virus is a DNA virus such as hepatitis B virus and the like. However, this publication does not include the compound disclosed in the present specification or a description regarding or suggestive of HCV.

Japanese Patent Application under PCT laid-open under kohyo No. 10-505092 discloses the following compound B and the like, wherein the use thereof is for a treatment of viral infection. The target virus is a DNA virus such as herpesvirus and hepatitis B virus. However, this publication does not include the compound disclosed in the present specification or a description regarding or suggestive of HCV.



compound A



compound B

The benzimidazole derivatives having an antiviral activity have been disclosed in JP-A-3-31264, US3644382 and US3778504. In addition, WO98/37072 discloses, as a production inhibitor of tumor necrosis factor (TNF) and cyclic AMP, a benzimidazole



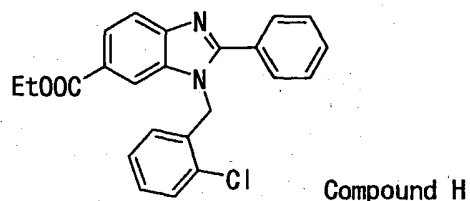
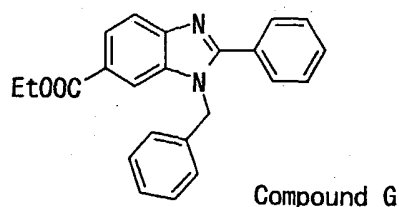
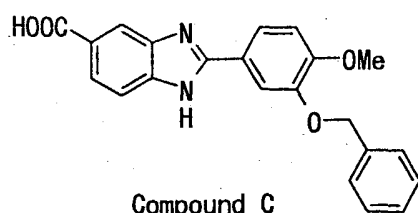
derivative for the use as an anti-human immunodeficiency virus (HIV) agent and an anti-inflammation agent. WO98/05327 discloses, as a reverse transcriptase inhibitor, a benzimidazole derivative for the use as an anti-HIV agent. J. Med. Chem. (13(4), 697-704, 5 1970) discloses, as a neuraminidase inhibitor, a benzimidazole derivative for the use as an anti-influenza virus agent.

However, none of these publications includes the compound of the present invention or a description regarding or suggestive of an anti-HCV effect.

10 Known benzimidazole derivatives having a pharmaceutical use other than as an antiviral agent are disclosed in JP-A-8-501318 (U.S. Patent 5,814,651) and JP-A-8-134073 (U.S. Patent 5,563,143). These publications disclose the following compound C and the like as a catechol diether compound, and the use thereof 15 as an anti-inflammation agent. However, neither of the publications includes the compound of the present invention, and as the action mechanism, the former discloses phosphodiesterase IV and the latter discloses TNF. These publications do not include a description regarding or suggestive of an anti-HCV 20 effect.

Japanese Patent Application under PCT laid-open under kohyo No. 2000-159749 (EP882718) discloses the following compound G and the like, and the use thereof for the treatment of bronchitis, glomerulonephritis and the like. However, this 25 publication does not include the compound of the present invention, but discloses only a phosphodiesterase IV inhibitory and hypoglycemic action. This publication does not include a description regarding or suggestive of an anti-HCV effect.

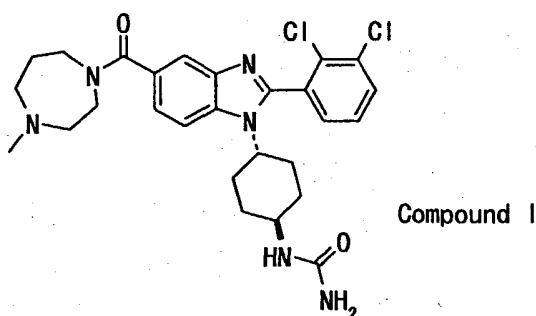
U.S. Patent 6,211,177 discloses the following compound H 30 and the like with their use as antitumor agents. However, this publication does not encompass the compound of the present invention, and does not disclose or suggest an anti-HCV effect.



WO98/50029, WO98/50030 and WO98/50031 disclose benzimidazole derivatives as an antitumor agent having a protein isoprenyl transferase action. While this publication discloses a wide scope of the claims, at least it does not include a compound analogous to the compound of the present invention or a description regarding or suggestive of an anti-HCV effect.

JP-A-8-109169 (EP694535) discloses the application of a tachykinin receptor antagonist to treat an inflammatory disease, and WO96/35713 discloses the application thereof as a growth hormone release promoter to treat a growth hormone-related disease such as osteoporosis and the like. However, none of these publications includes a description regarding or suggestive of an anti-HCV effect.

WO2001/21634 discloses the following compound I in a chemical library. However, this publication does not encompass the compound of the present invention. While it discloses an antimicrobial activity of certain compounds, this publication does not teach or suggest an anti-HCV effect.



JP-A-53-14735 discloses a benzimidazole derivative as a brightener besides its pharmaceutical use, but this publication does not include the compound of the present invention.

### Summary of the Invention

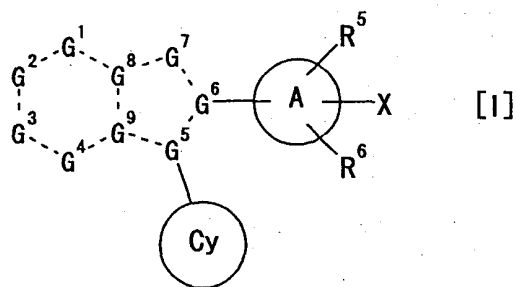
5 Based on the findings from the preceding studies, it has been elucidated that a pharmaceutical agent having an anti-HCV activity is effective for the prophylaxis and treatment of hepatitis C, and particularly an anti-HCV agent having an inhibitory activity on RNA-dependent RNA polymerase of HCV can be  
10 a prophylactic and therapeutic agent effective against hepatitis C and a prophylactic and therapeutic agent for the disease caused by hepatitis C.

Accordingly, the present invention provides a pharmaceutical agent having an anti-HCV activity, particularly a  
15 pharmaceutical agent having an RNA-dependent RNA polymerase inhibitory activity.

The present inventors have made an in-depth study of compounds having an anti-HCV activity, particularly RNA-dependent RNA polymerase inhibitory activity, and completed the present  
20 invention.

Thus, the present invention provides the following (1) to (117).

(1) A therapeutic agent for hepatitis C, which comprises a fused ring compound of the following formula [I] or a pharmaceutically  
25 acceptable salt thereof as an active ingredient:



wherein

a broken line is a single bond or a double bond,

G<sup>1</sup> is C(-R<sup>1</sup>) or a nitrogen atom,

30 G<sup>2</sup> is C(-R<sup>2</sup>) or a nitrogen atom,

G<sup>3</sup> is C(-R<sup>3</sup>) or a nitrogen atom,

G<sup>4</sup> is C(-R<sup>4</sup>) or a nitrogen atom,

$G^5$ ,  $G^6$ ,  $G^8$  and  $G^9$  are each independently a carbon atom or a nitrogen atom,  
 $G^7$  is  $C(-R^7)$ , an oxygen atom, a sulfur atom, or a nitrogen atom optionally substituted by  $R^8$ ,  
 wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently,  
 (1) hydrogen atom,  
 (2)  $C_{1-6}$  alkanoyl,  
 (3) carboxyl,  
 (4) cyano,  
 (5) nitro,  
 (6)  $C_{1-6}$  alkyl optionally substituted by 1 to 3 substituent(s) selected from the following group A,  
 group A; halogen atom, hydroxyl group, carboxyl, amino,  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkoxy  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkoxycarbonyl and  $C_{1-6}$  alkylamino,  
 (7)  $-COOR^{a1}$   
 wherein  $R^{a1}$  is optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the following group B or glucuronic acid residue,  
 group B; halogen atom, cyano, nitro,  $C_{1-6}$  alkyl, halogenated  $C_{1-6}$  alkyl,  $C_{1-6}$  alkanoyl,  
 $-(CH_2)_r-COOR^{b1}$ ,  $-(CH_2)_r-CONR^{b1}R^{b2}$ ,  $-(CH_2)_r-NR^{b1}R^{b2}$ ,  
 $-(CH_2)_r-NR^{b1}-COR^{b2}$ ,  $-(CH_2)_r-NHSO_2R^{b1}$ ,  $-(CH_2)_r-OR^{b1}$ ,  
 $-(CH_2)_r-SR^{b1}$ ,  $-(CH_2)_r-SO_2R^{b1}$  and  $-(CH_2)_r-SO_2NR^{b1}R^{b2}$   
 wherein  $R^{b1}$  and  $R^{b2}$  are each independently hydrogen atom or  $C_{1-6}$  alkyl and  $r$  is 0 or an integer of 1 to 6,  
 (8)  $-CONR^{a2}R^{a3}$   
 wherein  $R^{a2}$  and  $R^{a3}$  are each independently hydrogen atom,  $C_{1-6}$  alkoxy or optionally substituted  $C_{1-6}$  alkyl (as defined above),  
 (9)  $-C(=NR^{a4})NH_2$   
 wherein  $R^{a4}$  is hydrogen atom or hydroxyl group,  
 (10)  $-NHR^{a5}$   
 wherein  $R^{a5}$  is hydrogen atom,  $C_{1-6}$  alkanoyl or  $C_{1-6}$  alkylsulfonyl,  
 (11)  $-OR^{a6}$

wherein  $R^{a6}$  is hydrogen atom or optionally substituted  $C_{1-6}$  alkyl(as defined above),

(12)  $-SO_2R^{a7}$

wherein  $R^{a7}$  is hydroxyl group, amino,  $C_{1-6}$  alkyl or  $C_{1-6}$  alkylamino,

(13)  $-P(=O)(OR^{a31})_2$

wherein  $R^{a31}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B

or

(14) heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom, and

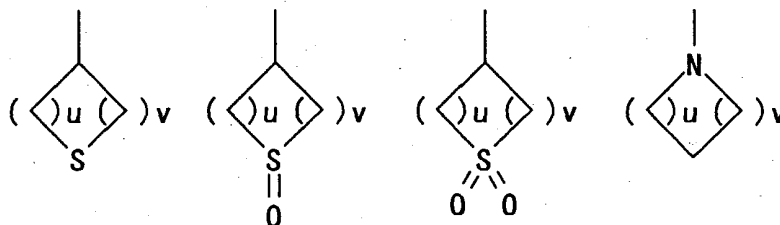
$R^7$  and  $R^8$  are each hydrogen atom or optionally substituted  $C_{1-6}$  alkyl (as defined above),

ring Cy is

(1)  $C_{3-8}$  cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the following group C, group C; hydroxyl group, halogen atom,  $C_{1-6}$  alkyl and  $C_{1-6}$  alkoxy,

(2)  $C_{3-8}$  cycloalkenyl optionally substituted by 1 to 5 substituent(s) selected from the above group C, or

(3)



wherein u and v are each independently an integer of 1 to 3,

ring A is

(1)  $C_{6-14}$  aryl,

(2)  $C_{3-8}$  cycloalkyl,

(3)  $C_{3-8}$  cycloalkenyl or

(4) heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom,

R<sup>5</sup> and R<sup>6</sup> are each independently

- (1) hydrogen atom,
- (2) halogen atom,
- (3) optionally substituted C<sub>1-6</sub> alkyl (as defined above)

5 or

- (4) -OR<sup>a8</sup>

wherein R<sup>a8</sup> is hydrogen atom, C<sub>1-6</sub> alkyl or C<sub>6-14</sub> aryl  
C<sub>1-6</sub> alkyl, and

X is

- 10 (1) hydrogen atom,
- (2) halogen atom,
- (3) cyano,
- (4) nitro,
- (5) amino, C<sub>1-6</sub> alkanoylamino,
- 15 (6) C<sub>1-6</sub> alkylsulfonyl,
- (7) optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- (8) C<sub>2-6</sub> alkenyl optionally substituted by 1 to 3  
substituent(s) selected from the above group A,
- (9) -COOR<sup>a9</sup>

20

wherein R<sup>a9</sup> is hydrogen atom or C<sub>1-6</sub> alkyl,

- (10) -CONH-(CH<sub>2</sub>)<sub>1</sub>-R<sup>a10</sup>

wherein R<sup>a10</sup> is optionally substituted C<sub>1-6</sub> alkyl (as  
defined above), C<sub>1-6</sub> alkoxy carbonyl or C<sub>1-6</sub>  
alkanoylamino and 1 is 0 or an integer of 1 to 6,

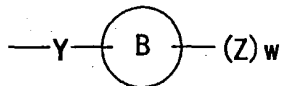
25

- (11) -OR<sup>a11</sup>

wherein R<sup>a11</sup> is hydrogen atom or optionally substituted  
C<sub>1-6</sub> alkyl (as defined above)

or

- (12)



30

wherein

ring B is

- (1') C<sub>6-14</sub> aryl,
- (2') C<sub>3-8</sub> cycloalkyl or
- 35 (3') heterocyclic group (as defined above),

each Z is independently

- (1') a group selected from the following group D,

(2') C<sub>6-14</sub> aryl optionally substituted by 1 to 5  
substituent(s) selected from the following group  
D,

(3') C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to  
5 substituent(s) selected from the following  
group D,

(4') C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by  
1 to 5 substituent(s) selected from the  
following group D,

(5') heterocyclic group optionally substituted by 1  
to 5 substituent(s) selected from the  
following group D,

wherein the heterocyclic group has 1 to 4 hetero-  
atom(s) selected from an oxygen atom, a nitrogen  
atom and a sulfur atom, or

(6') heterocycle C<sub>1-6</sub> alkyl optionally substituted  
by 1 to 5 substituent(s) selected from the  
following group D,

wherein the heterocycle C<sub>1-6</sub> alkyl is C<sub>1-6</sub> alkyl  
substituted by heterocyclic group optionally  
substituted by 1 to 5 substituent(s) selected from  
the group D, as defined above,

group D:

(a) hydrogen atom,

(b) halogen atom,

(c) cyano,

(d) nitro,

(e) optionally substituted C<sub>1-6</sub> alkyl (as defined  
above),

(f) -(CH<sub>2</sub>)<sub>t</sub>-COR<sup>a18</sup>,

(hereinafter each t means independently 0 or an  
integer of 1 to 6),

wherein R<sup>a18</sup> is

(1") optionally substituted C<sub>1-6</sub> alkyl (as  
defined above),

(2") C<sub>6-14</sub> aryl optionally substituted by 1 to  
5 substituent(s) selected from the above  
group B or

- (3") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B wherein the heterocyclic group has 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom,
- 5
- (g)  $-(CH_2)_t-COOR^{a19}$  wherein  $R^{a19}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 10
- (h)  $-(CH_2)_t-CONR^{a27}R^{a28}$  wherein  $R^{a27}$  and  $R^{a28}$  are each independently,
- 15
- (1") hydrogen atom,
- (2") optionally substituted  $C_{1-6}$  alkyl (as defined above),
- (3")  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 20
- (4")  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (5") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 25
- (6") heterocycle  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 30
- wherein the heterocycle  $C_{1-6}$  alkyl is  $C_{1-6}$  alkyl substituted by heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B, as defined above,
- (7")  $C_{3-8}$  cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 35
- (8")  $C_{3-8}$  cycloalkyl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected



from the above group B,

(9") hydroxyl group or

(10") C<sub>1-6</sub> alkoxy,

(i)  $-(CH_2)_t-C(=NR^{a33})NH_2$

5 wherein R<sup>a33</sup> is hydrogen atom, C<sub>1-6</sub> alkyl,  
hydroxyl group or C<sub>1-6</sub> alkoxy,

(j)  $-(CH_2)_t-OR^{a20}$

wherein R<sup>a20</sup> is

(1") hydrogen atom,

10 (2") optionally substituted C<sub>1-6</sub> alkyl (as  
defined above),

(3") optionally substituted C<sub>2-6</sub> alkenyl (as  
defined above),

15 (4") C<sub>2-6</sub> alkynyl optionally substituted by 1  
to 3 substituent(s) selected from the  
above group A,

(5") C<sub>6-14</sub> aryl optionally substituted by 1 to  
5 substituent(s) selected from the  
above group B,

20 (6") C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

(7") heterocyclic group optionally  
substituted by 1 to 5 substituent(s)  
25 selected from the above group B,

(8") heterocycle C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

30 (9") C<sub>3-8</sub> cycloalkyl optionally substituted by  
1 to 5 substituent(s) selected from the  
above group B, or

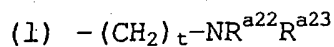
(10") C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

35 (k)  $-(CH_2)_t-O-(CH_2)_p-COR^{a21}$

wherein R<sup>a21</sup> is amino, C<sub>1-6</sub> alkylamino or  
heterocyclic group optionally substituted by

1 to 5 substituent(s) selected from the above group B,

and p is 0 or an integer of 1 to 6,



wherein  $R^{a22}$  and  $R^{a23}$  are each independently

(1") hydrogen atom,

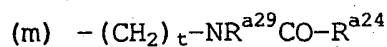
(2") optionally substituted  $C_{1-6}$  alkyl (as defined above),

(3")  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4")  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(5") heterocycle  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B or

(6") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,



wherein  $R^{a29}$  is hydrogen atom,  $C_{1-6}$  alkyl or  $C_{1-6}$  alkanoyl, and

$R^{a24}$  is

(1") amino,

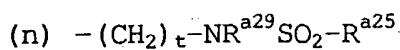
(2")  $C_{1-6}$  alkylamino,

(3") optionally substituted  $C_{1-6}$  alkyl (as defined above),

(4")  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(5") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B or

(6") heterocycle  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,



wherein  $R^{a29}$  is as defined above, and  
 $R^{a25}$  is hydrogen atom, optionally  
substituted  $C_{1-6}$  alkyl (as defined above),  
 $C_{6-14}$  aryl optionally substituted by 1 to 5  
substituent(s) selected from the above group  
B or heterocyclic group optionally  
substituted by 1 to 5 substituent(s) selected  
from the above group B,

(o)  $-(CH_2)_t-S(O)_q-R^{a25}$

wherein  $R^{a25}$  is as defined above, and  $q$  is 0,  
1 or 2,

(p)  $-(CH_2)_t-SO_2-NHR^{a26}$

wherein  $R^{a26}$  is hydrogen atom, optionally  
substituted  $C_{1-6}$  alkyl (as defined above),  
 $C_{6-14}$  aryl optionally substituted by 1 to 5  
substituent(s) selected from the above group  
B or heterocyclic group optionally  
substituted by 1 to 5 substituent(s) selected  
from the above group B,

and

(q) heterocyclic group having 1 to 4

heteroatom(s) selected from an oxygen atom,  
a nitrogen atom and a sulfur atom, and

$w$  is an integer of 1 to 3, and

$Y$  is

(1') a single bond,

(2')  $C_{1-6}$  alkylene,

(3')  $C_{2-6}$  alkenylene,

(4')  $-(CH_2)_m-O-(CH_2)_n-$ ,

(hereinafter  $m$  and  $n$  are each independently 0  
or an integer of 1 to 6),

(5')  $-CO-$ ,

(6')  $-CO_2-(CH_2)_n-$ ,

(7')  $-CONH-(CH_2)_n-NH-$ ,

(8')  $-NHCO_2-$ ,

(9')  $-NHCONH-$ ,

(10')  $-O-(CH_2)_n-CO-$ ,

(11')  $-O-(CH_2)_n-O-$ ,

(12')  $-\text{SO}_2-$ ,

(13')  $-(\text{CH}_2)_m-\text{NR}^{\text{a12}}-(\text{CH}_2)_n-$

wherein  $\text{R}^{\text{a12}}$  is

(1'') hydrogen atom,

(2'') optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

(3'')  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4'')  $\text{C}_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(5'')  $-\text{COR}^{\text{b5}}$

wherein  $\text{R}^{\text{b5}}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),  $\text{C}_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(6'')  $-\text{COOR}^{\text{b5}}$  ( $\text{R}^{\text{b5}}$  is as defined above) or

(7'')  $-\text{SO}_2\text{R}^{\text{b5}}$  ( $\text{R}^{\text{b5}}$  is as defined above),

(14')  $-\text{NR}^{\text{a12}}\text{CO}-$  ( $\text{R}^{\text{a12}}$  is as defined above),

(15')  $-\text{CONR}^{\text{a13}}-(\text{CH}_2)_n-$

wherein  $\text{R}^{\text{a13}}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above) or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(16')  $-\text{CONH}-\text{CHR}^{\text{a14}}-$

wherein  $\text{R}^{\text{a14}}$  is  $\text{C}_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(17')  $-\text{O}-(\text{CH}_2)_m-\text{CR}^{\text{a15}}\text{R}^{\text{a16}}-(\text{CH}_2)_n-$

wherein  $\text{R}^{\text{a15}}$  and  $\text{R}^{\text{a16}}$  are each independently

(1'') hydrogen atom,

(2'') carboxyl,

(3'') C<sub>1-6</sub> alkyl,

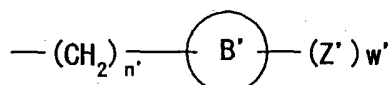
(4'') -OR<sup>b6</sup>

wherein R<sup>b6</sup> is C<sub>1-6</sub> alkyl or C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl, or

(5'') -NHR<sup>b7</sup>

wherein R<sup>b7</sup> is hydrogen atom, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkanoyl or C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyloxycarbonyl, or R<sup>a15</sup> is optionally

(6'')



wherein n', ring B', Z' and w' are the same as the above-mentioned n, ring B, Z and w, respectively, and may be the same as or different from the respective counterparts,

(18') -(CH<sub>2</sub>)<sub>n</sub>-NR<sup>a12</sup>-CHR<sup>a15</sup>- (R<sup>a12</sup> and R<sup>a15</sup> are each as defined above),

(19') -NR<sup>a17</sup>SO<sub>2</sub>-

wherein R<sup>a17</sup> is hydrogen atom or C<sub>1-6</sub> alkyl,

(20') -S(O)<sub>e</sub>-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>- (e is 0, 1 or 2, R<sup>a15</sup> and R<sup>a16</sup> are each as defined above),

or

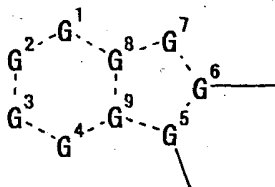
(21') -(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>- (R<sup>a15</sup> and R<sup>a16</sup> are each as defined above).

(2) The therapeutic agent of (1) above, wherein 1 to 4 of the G<sup>1</sup>, G<sup>2</sup>, G<sup>3</sup>, G<sup>4</sup>, G<sup>5</sup>, G<sup>6</sup>, G<sup>7</sup>, G<sup>8</sup> and G<sup>9</sup> is(are) a nitrogen atom.

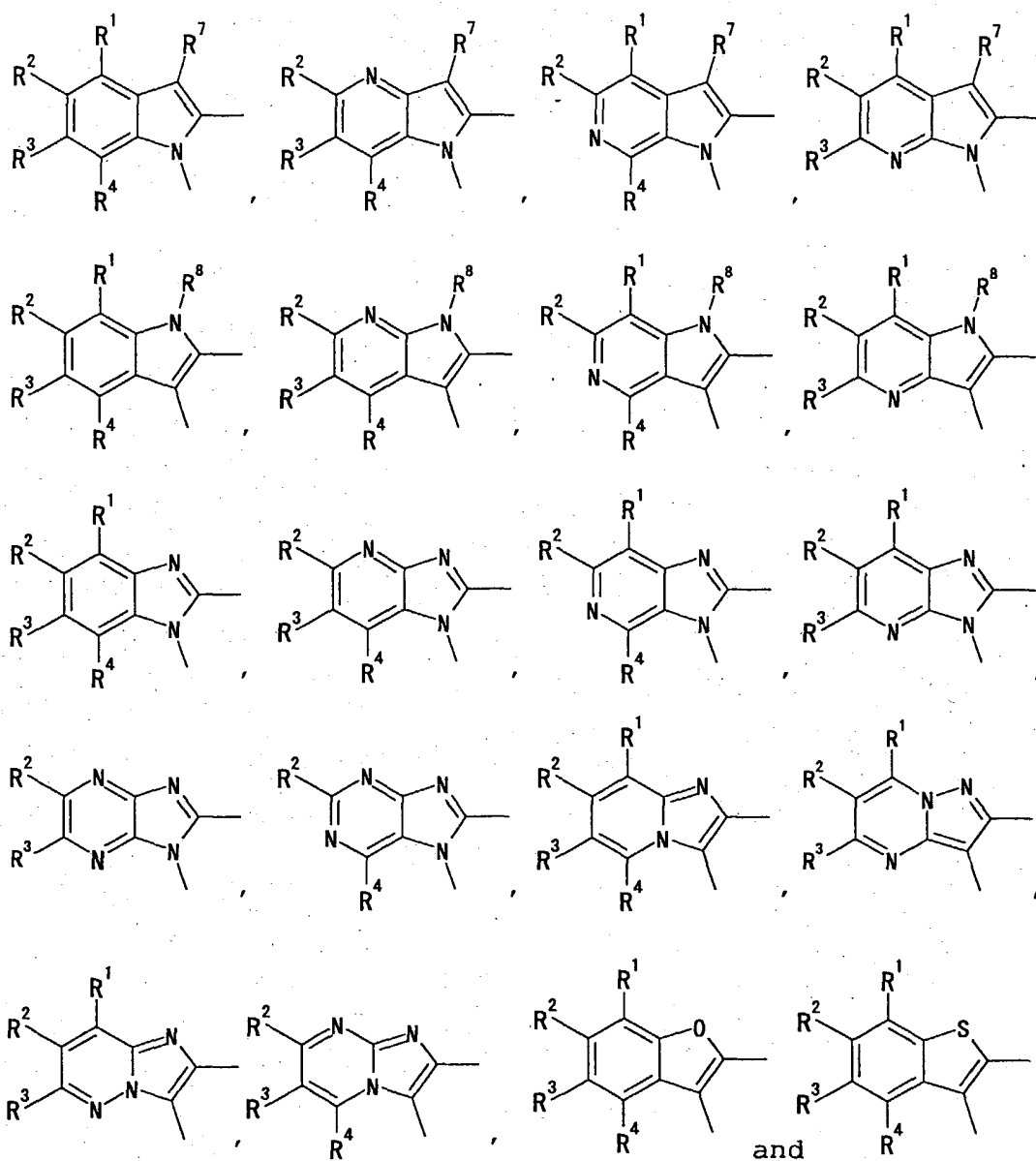
(3) The therapeutic agent of (2) above, wherein G<sup>2</sup> is C(-R<sup>2</sup>) and G<sup>6</sup> is a carbon atom.

(4) The therapeutic agent of (2) or (3) above, wherein G<sup>5</sup> is a nitrogen atom.

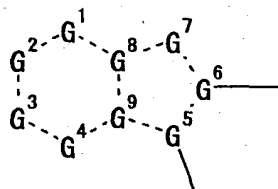
(5) The therapeutic agent of (1) above, wherein, in formula [I], the moiety



is a fused ring selected from

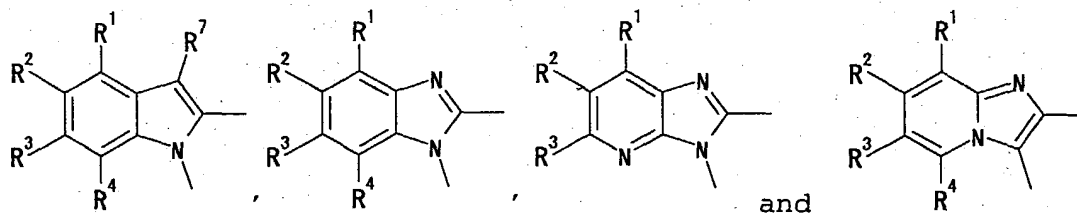


(6) The therapeutic agent of (5) above, wherein, in formula [I], the moiety

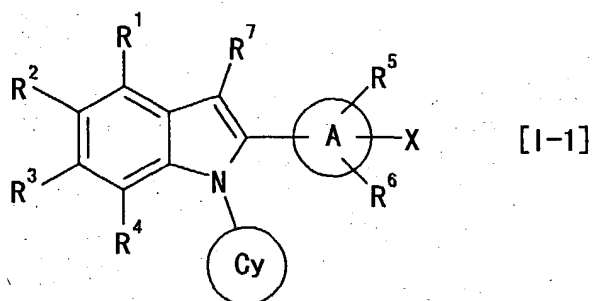


5

is a fused ring selected from

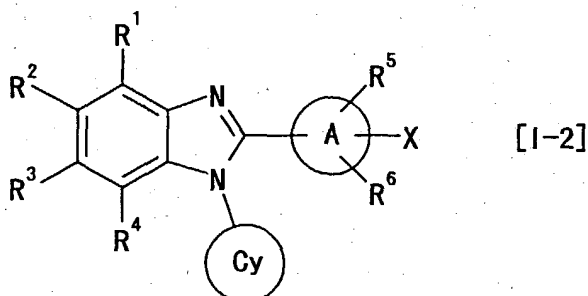


(7) The therapeutic agent of (6) above, which comprises a fused ring compound of the following formula [I-1]



5 wherein each symbol is as defined in (1),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

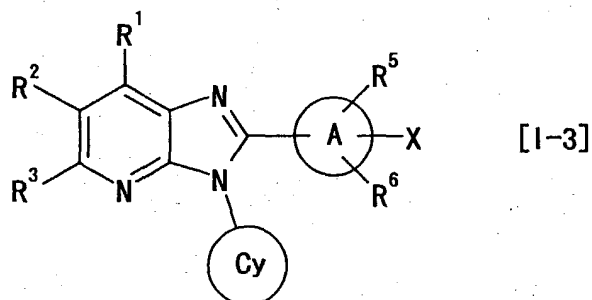
(8) The therapeutic agent of (6) above, which comprises a fused ring compound of the following formula [I-2]



10

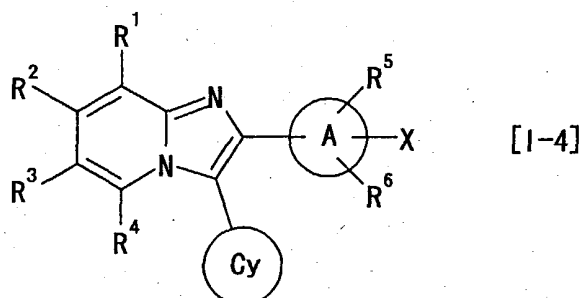
wherein each symbol is as defined in (1),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

(9) The therapeutic agent of (6) above, which comprises a fused  
15 ring compound of the following formula [I-3]



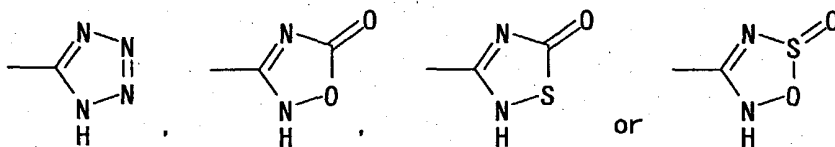
wherein each symbol is as defined in (1),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

(10) The therapeutic agent of (6) above, which comprises a fused  
5 ring compound of the following formula [I-4]



wherein each symbol is as defined in (1),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

10 (11) The therapeutic agent of any of (1) to (10) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl,  $-\text{COOR}^{a1}$ ,  $-\text{CONR}^{a2}\text{R}^{a3}$ ,  $-\text{SO}_2\text{R}^{a7}$  (wherein  $R^{a1}$ ,  $R^{a2}$ ,  $R^{a3}$  and  $R^{a7}$  are as defined in (1)),



(12) The therapeutic agent of (11) above, wherein at least one of  
15  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl,  $-\text{COOR}^{a1}$ ,  $-\text{CONR}^{a2}\text{R}^{a3}$  or  $-\text{SO}_2\text{R}^{a7}$  wherein  $R^{a1}$ ,  $R^{a2}$ ,  $R^{a3}$  and  $R^{a7}$  are as defined in (1).

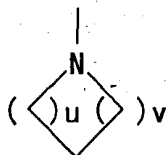
(13) The therapeutic agent of any of (1) to (10) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is  $-\text{COOR}^{a1}$  wherein  $R^{a1}$  is glucuronic acid residue.

20 (14) The therapeutic agent of any of (1) to (10) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom.

(15) The therapeutic agent of any of (1) to (14) above, wherein  
25 the ring Cy is cyclopentyl, cyclohexyl, cycloheptyl, tetrahydrothiopyranyl or piperidino.

(16) The therapeutic agent of any of (1) to (14) above, wherein the ring Cy is





wherein each symbol is as defined in (1).

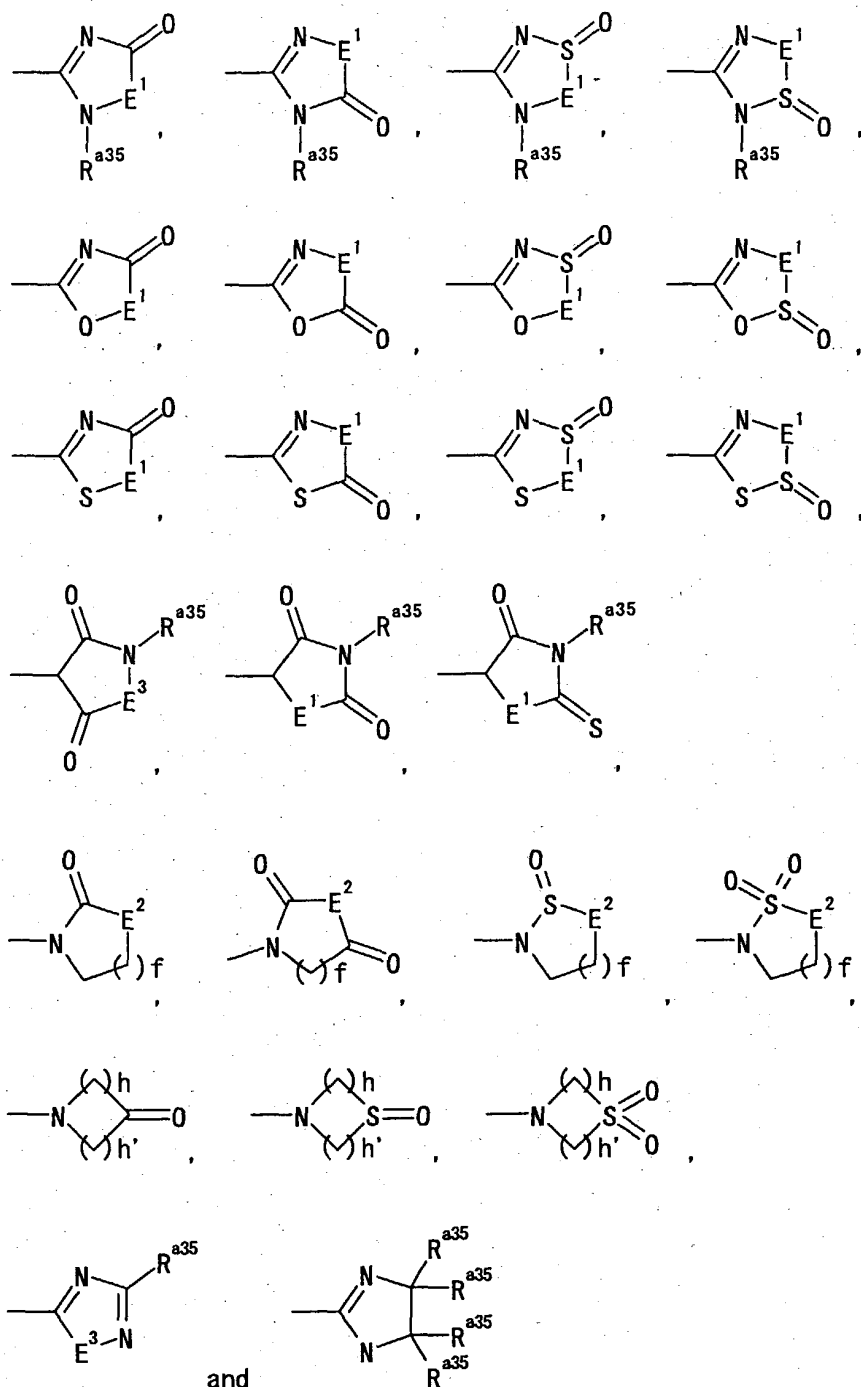
(17) The therapeutic agent of any of (1) to (16) above, wherein the ring A is C<sub>6-14</sub> aryl.

5 (18) The therapeutic agent of any of (1) to (17) above, wherein at least one substituent optionally substituted by group A is a substituent substituted by C<sub>1-6</sub> alkoxy C<sub>1-6</sub> alkoxy.

(19) The therapeutic agent of any of (1) to (17) above, wherein the Y is -(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>- wherein each symbol is as defined  
10 in (1).

(20) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by Z is heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the group D.

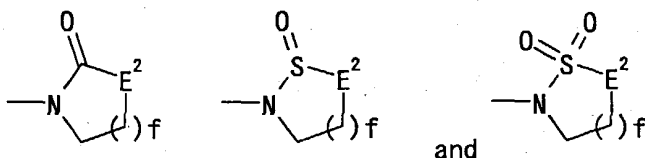
15 (21) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by Z is a heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the group D, wherein said heterocyclic group is selected from the following groups:



5 wherein  $E^1$  is an oxygen atom, a sulfur atom or  $N(-R^{a35})$ ,  $E^2$  is an oxygen atom,  $CH_2$  or  $N(-R^{a35})$ ,  $E^3$  is an oxygen atom or a sulfur atom, wherein each  $R^{a35}$  is independently hydrogen atom or  $C_{1-6}$  alkyl,  $f$  is an integer of 1 to 3, and  $h$  and  $h'$  are the same or different and each is an integer of 1 to 3.

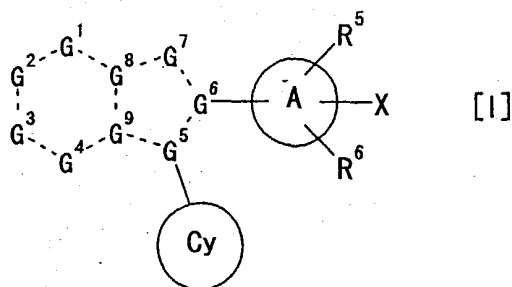
10 (22) The therapeutic agent of (21) above, wherein at least one group represented by  $Z$  is heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the group D

wherein said heterocyclic group is selected from the following groups:



wherein each symbol is as defined in (21).

- 5 (23) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is  $-(CH_2)_t-CONR^{a27}R^{a28}$  wherein each symbol is as defined in (1), and at least one of  $R^{a27}$  and  $R^{a28}$  is  $C_{1-6}$  alkoxy.
- (24) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is  $-(CH_2)_t-C(=NR^{a33})NH_2$  wherein each symbol is as defined in (1), and  $R^{a33}$  is hydroxyl group or  $C_{1-6}$  alkoxy.
- (25) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is  $-(CH_2)_t-O-(CH_2)_p-COR^{a21}$  15 wherein each symbol is as defined in (1), and  $R^{a21}$  is amino.
- (26) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is  $-(CH_2)_t-NR^{a29}CO-R^{a24}$  wherein each symbol is as defined in (1), and  $R^{a24}$  is amino or  $C_{1-6}$  alkylamino.
- 20 (27) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is  $-(CH_2)_t-NR^{a22}R^{a23}$  wherein each symbol is as defined in (1), and at least one of  $R^{a22}$  and  $R^{a23}$  is heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the group B.
- 25 (28) The therapeutic agent of any of (1) to (19) above, wherein at least one group represented by group D is heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom.
- (29) The therapeutic agent of (1) above, which comprises a fused 30 ring compound of the following formula [I] or a pharmaceutically acceptable salt thereof as an active ingredient:



wherein

a broken line is a single bond or a double bond,

$G^1$  is  $C(-R^1)$  or a nitrogen atom,

5  $G^2$  is  $C(-R^2)$  or a nitrogen atom,

$G^3$  is  $C(-R^3)$  or a nitrogen atom,

$G^4$  is  $C(-R^4)$  or a nitrogen atom,

$G^5$ ,  $G^6$ ,  $G^8$  and  $G^9$  are each independently a carbon atom or a nitrogen atom,

10  $G^7$  is  $C(-R^7)$ , an oxygen atom, a sulfur atom, or a nitrogen atom optionally substituted by  $R^8$ ,

wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently,

(1) hydrogen atom,

(2)  $C_{1-6}$  alkanoyl,

15 (3) carboxyl,

(4) cyano,

(5) nitro,

(6)  $C_{1-6}$  alkyl optionally substituted by 1 to 3

substituent(s) selected from the following group A,

20 group A; halogen atom, hydroxyl group, carboxyl, amino,  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkoxy carbonyl and  $C_{1-6}$  alkylamino,

(7)  $-COOR^{a1}$

wherein  $R^{a1}$  is optionally substituted  $C_{1-6}$  alkyl (as

25 defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally

substituted by 1 to 5 substituent(s) selected from the following group B,

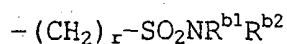
group B; halogen atom, cyano, nitro,  $C_{1-6}$  alkyl,

halogenated  $C_{1-6}$  alkyl,  $C_{1-6}$  alkanoyl,

30  $-(CH_2)_r-COOR^{b1}$ ,  $-(CH_2)_r-CONR^{b1}R^{b2}$ ,  $-(CH_2)_r-NR^{b1}R^{b2}$ ,

$-(CH_2)_r-NR^{b1}-COR^{b2}$ ,  $-(CH_2)_r-NHSO_2R^{b1}$ ,

$-(CH_2)_r-OR^{b1}$ ,  $-(CH_2)_r-SR^{b1}$ ,  $-(CH_2)_r-SO_2R^{b1}$  and



wherein  $\text{R}^{b1}$  and  $\text{R}^{b2}$  are each independently hydrogen atom or  $\text{C}_{1-6}$  alkyl and  $r$  is 0 or an integer of 1 to 6,

5 (8)  $-\text{CONR}^{a2}\text{R}^{a3}$

wherein  $\text{R}^{a2}$  and  $\text{R}^{a3}$  are each independently hydrogen atom,  $\text{C}_{1-6}$  alkoxy or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

(9)  $-\text{C}(=\text{NR}^{a4})\text{NH}_2$

10 wherein  $\text{R}^{a4}$  is hydrogen atom or hydroxyl group,

(10)  $-\text{NHR}^{a5}$

wherein  $\text{R}^{a5}$  is hydrogen atom,  $\text{C}_{1-6}$  alkanoyl or  $\text{C}_{1-6}$  alkylsulfonyl,

(11)  $-\text{OR}^{a6}$

15 wherein  $\text{R}^{a6}$  is hydrogen atom or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

(12)  $-\text{SO}_2\text{R}^{a7}$

wherein  $\text{R}^{a7}$  is hydroxyl group, amino,  $\text{C}_{1-6}$  alkyl or  $\text{C}_{1-6}$  alkylamino

20 or

(13)  $-\text{P}(=\text{O})(\text{OR}^{a31})_2$

wherein  $\text{R}^{a31}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above) or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s)

25 selected from the above group B, and

$\text{R}^7$  and  $\text{R}^8$  are each hydrogen atom or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

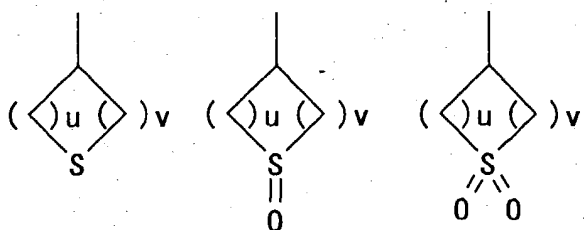
ring Cy is

(1)  $\text{C}_{3-8}$  cycloalkyl optionally substituted by 1 to 5

30 substituent(s) selected from the following group C, group C; hydroxyl group, halogen atom,  $\text{C}_{1-6}$  alkyl and  $\text{C}_{1-6}$  alkoxy,

(2)  $\text{C}_{3-8}$  cycloalkenyl optionally substituted by 1 to 5 substituent(s) selected from the above group C, or

35 (3)



wherein u and v are each independently an integer of 1 to 3,

ring A is

- (1) C<sub>6-14</sub> aryl,
- (2) C<sub>3-8</sub> cycloalkyl,
- (3) C<sub>3-8</sub> cycloalkenyl or
- (4) heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom,

R<sup>5</sup> and R<sup>6</sup> are each independently

- (1) hydrogen atom,
- (2) halogen atom,
- (3) optionally substituted C<sub>1-6</sub> alkyl (as defined above) or
- (4) -OR<sup>a8</sup>

wherein R<sup>a8</sup> is hydrogen atom, C<sub>1-6</sub> alkyl or C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl, and

X is

- (1) hydrogen atom,
- (2) halogen atom,
- (3) cyano,
- (4) nitro,
- (5) amino, C<sub>1-6</sub> alkanoylamino,
- (6) C<sub>1-6</sub> alkylsulfonyl,
- (7) optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- (8) C<sub>2-6</sub> alkenyl optionally substituted by 1 to 3 substituent(s) selected from the above group A,
- (9) -COOR<sup>a9</sup>

wherein R<sup>a9</sup> is hydrogen atom or C<sub>1-6</sub> alkyl,

- (10) -CONH-(CH<sub>2</sub>)<sub>1</sub>-R<sup>a10</sup>

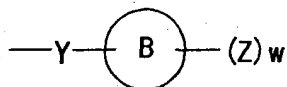
wherein R<sup>a10</sup> is optionally substituted C<sub>1-6</sub> alkyl (as defined above), C<sub>1-6</sub> alkoxy carbonyl or C<sub>1-6</sub>

alkanoylamino and 1 is 0 or an integer of 1 to 6,  
(11)  $-OR^{a11}$

wherein  $R^{a11}$  is hydrogen atom or optionally substituted  
 $C_{1-6}$  alkyl (as defined above)

5 or

(12)



wherein

ring B is

10 (1')  $C_{6-14}$  aryl,

(2')  $C_{3-8}$  cycloalkyl or

(3') heterocyclic group (as defined above),

each Z is independently

(1') a group selected from the following group D,

15 (2')  $C_{6-14}$  aryl optionally substituted by 1 to 5  
substituent(s) selected from the following group  
D,

(3')  $C_{3-8}$  cycloalkyl optionally substituted by 1 to 5  
substituent(s) selected from the following group  
20 D,

(4')  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by  
1 to 5 substituent(s) selected from the  
following group D or

25 (5') heterocyclic group optionally substituted by 1  
to 5 substituent(s) selected from the  
following group D

wherein the heterocyclic group has 1 to 4 hetero-  
atom(s) selected from an oxygen atom, a nitrogen  
atom and a sulfur atom,

30 group D:

(a) hydrogen atom,

(b) halogen atom,

(c) cyano,

(d) nitro,

35 (e) optionally substituted  $C_{1-6}$  alkyl (as defined  
above),

(f)  $-(CH_2)_t-COR^{a18}$ ,

(hereinafter each t means independently 0 or an integer of 1 to 6),

wherein R<sup>a18</sup> is

(1") optionally substituted C<sub>1-6</sub> alkyl (as defined above),

(2") C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or

(3") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B wherein the heterocyclic group has 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom,

(g) -(CH<sub>2</sub>)<sub>t</sub>-COOR<sup>a19</sup>

wherein R<sup>a19</sup> is hydrogen atom, optionally substituted C<sub>1-6</sub> alkyl (as defined above) or C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(h) -(CH<sub>2</sub>)<sub>t</sub>-CONR<sup>a27</sup>R<sup>a28</sup>

wherein R<sup>a27</sup> and R<sup>a28</sup> are each independently,

(1") hydrogen atom,

(2") optionally substituted C<sub>1-6</sub> alkyl (as defined above),

(3") C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4") C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(5") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(6") heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B, wherein the heterocycle C<sub>1-6</sub> alkyl is C<sub>1-6</sub>



alkyl substituted by heterocyclic group  
optionally substituted by 1 to 5 substituent(s)  
selected from the above group B, as defined  
above,

5 (7") C<sub>3-8</sub> cycloalkyl optionally substituted by 1  
to 5 substituent(s) selected from the above  
group B, or

(8") C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s) selected  
10 from the above group B,

(i)  $-(CH_2)_t-C(=NR^{a33})NH_2$   
wherein R<sup>a33</sup> is hydrogen atom or C<sub>1-6</sub> alkyl,

(j)  $-(CH_2)_t-OR^{a20}$   
wherein R<sup>a20</sup> is

15 (1") hydrogen atom,

(2") optionally substituted C<sub>1-6</sub> alkyl (as  
defined above),

(3") optionally substituted C<sub>2-6</sub> alkenyl (as  
defined above),

20 (4") C<sub>2-6</sub> alkynyl optionally substituted by 1  
to 3 substituent(s) selected from the  
above group A,

(5") C<sub>6-14</sub> aryl optionally substituted by 1 to  
5 substituent(s) selected from the  
25 above group B,

(6") C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

(7") heterocyclic group optionally  
substituted by 1 to 5 substituent(s)  
30 selected from the above group B,

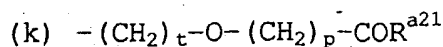
(8") heterocycle C<sub>1-6</sub> alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

35 (9") C<sub>3-8</sub> cycloalkyl optionally substituted by  
1 to 5 substituent(s) selected from the  
above group B, or

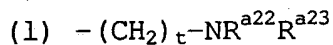
(10") C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally

substituted by 1 to 5 substituent(s)

selected from the above group B,



wherein  $R^{a21}$  is  $C_{1-6}$  alkylamino or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B, and p is 0 or an integer of 1 to 6,



wherein  $R^{a22}$  and  $R^{a23}$  are each independently

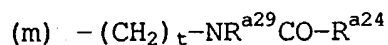
(1") hydrogen atom,

(2") optionally substituted  $C_{1-6}$  alkyl (as defined above),

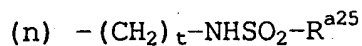
(3")  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4")  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B or

(5") heterocycle  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,



wherein  $R^{a29}$  is hydrogen atom,  $C_{1-6}$  alkyl or  $C_{1-6}$  alkanoyl,  $R^{a24}$  is optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,



wherein  $R^{a25}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(o)  $-(CH_2)_t-S(O)_q-R^{a25}$

wherein  $R^{a25}$  is as defined above, and q is 0, 1 or 2,

and

(p)  $-(CH_2)_t-SO_2-NHR^{a26}$

wherein  $R^{a26}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

w is an integer of 1 to 3, and

Y is

(1') a single bond,

(2')  $C_{1-6}$  alkylene,

(3')  $C_{2-6}$  alkenylene,

(4')  $-(CH_2)_m-O-(CH_2)_n-$ ,

(hereinafter m and n are each independently 0 or an integer of 1 to 6),

(5')  $-CO-$ ,

(6')  $-CO_2-(CH_2)_n-$ ,

(7')  $-CONH-(CH_2)_n-NH-$ ,

(8')  $-NHCO_2-$ ,

(9')  $-NHCONH-$ ,

(10')  $-O-(CH_2)_n-CO-$ ,

(11')  $-O-(CH_2)_n-O-$ ,

(12')  $-SO_2-$ ,

(13')  $-(CH_2)_m-NR^{a12}-(CH_2)_n-$

wherein  $R^{a12}$  is

(1'') hydrogen atom,

(2'') optionally substituted  $C_{1-6}$  alkyl (as defined above),

(3'')  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4'')  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the

above group B,

(5'')  $-\text{COR}^{\text{b5}}$

wherein  $\text{R}^{\text{b5}}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),  $\text{C}_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(6'')  $-\text{COOR}^{\text{b5}}$  ( $\text{R}^{\text{b5}}$  is as defined above) or

(7'')  $-\text{SO}_2\text{R}^{\text{b5}}$  ( $\text{R}^{\text{b5}}$  is as defined above),

(14')  $-\text{NR}^{\text{a12}}\text{CO}-$  ( $\text{R}^{\text{a12}}$  is as defined above),

(15')  $-\text{CONR}^{\text{a13}}-(\text{CH}_2)_n-$

wherein  $\text{R}^{\text{a13}}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above) or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(16')  $-\text{CONH}-\text{CHR}^{\text{a14}}-$

wherein  $\text{R}^{\text{a14}}$  is  $\text{C}_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(17')  $-\text{O}-(\text{CH}_2)_m-\text{CR}^{\text{a15}}\text{R}^{\text{a16}}-(\text{CH}_2)_n-$

wherein  $\text{R}^{\text{a15}}$  and  $\text{R}^{\text{a16}}$  are each independently

(1'') hydrogen atom,

(2'') carboxyl,

(3'')  $\text{C}_{1-6}$  alkyl,

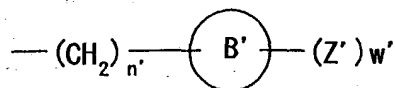
(4'')  $-\text{OR}^{\text{b6}}$

wherein  $\text{R}^{\text{b6}}$  is  $\text{C}_{1-6}$  alkyl or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl, or

(5'')  $-\text{NHR}^{\text{b7}}$

wherein  $\text{R}^{\text{b7}}$  is hydrogen atom,  $\text{C}_{1-6}$  alkyl,  $\text{C}_{1-6}$  alkanoyl or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyloxycarbonyl, or  $\text{R}^{\text{a15}}$  is optionally

(6'')



wherein  $n'$ , ring  $B'$ ,  $Z'$  and  $w'$  are the same as the above-mentioned  $n$ , ring  $B$ ,  $Z$  and  $w$ , respectively, and may be the same as or different from the respective counterparts,

(18')  $-(CH_2)_n-NR^{a12}-CHR^{a15}-$  ( $R^{a12}$  and  $R^{a15}$  are each as defined above),

(19')  $-NR^{a17}SO_2-$

wherein  $R^{a17}$  is hydrogen atom or  $C_{1-6}$  alkyl

or

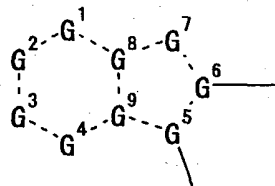
(20')  $-S(O)_e-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  ( $e$  is 0, 1 or 2,  $R^{a15}$  and  $R^{a16}$  are each as defined above).

(30) The therapeutic agent of (29) above, wherein 1 to 4 of the  $G^1$ ,  $G^2$ ,  $G^3$ ,  $G^4$ ,  $G^5$ ,  $G^6$ ,  $G^7$ ,  $G^8$  and  $G^9$  is(are) a nitrogen atom.

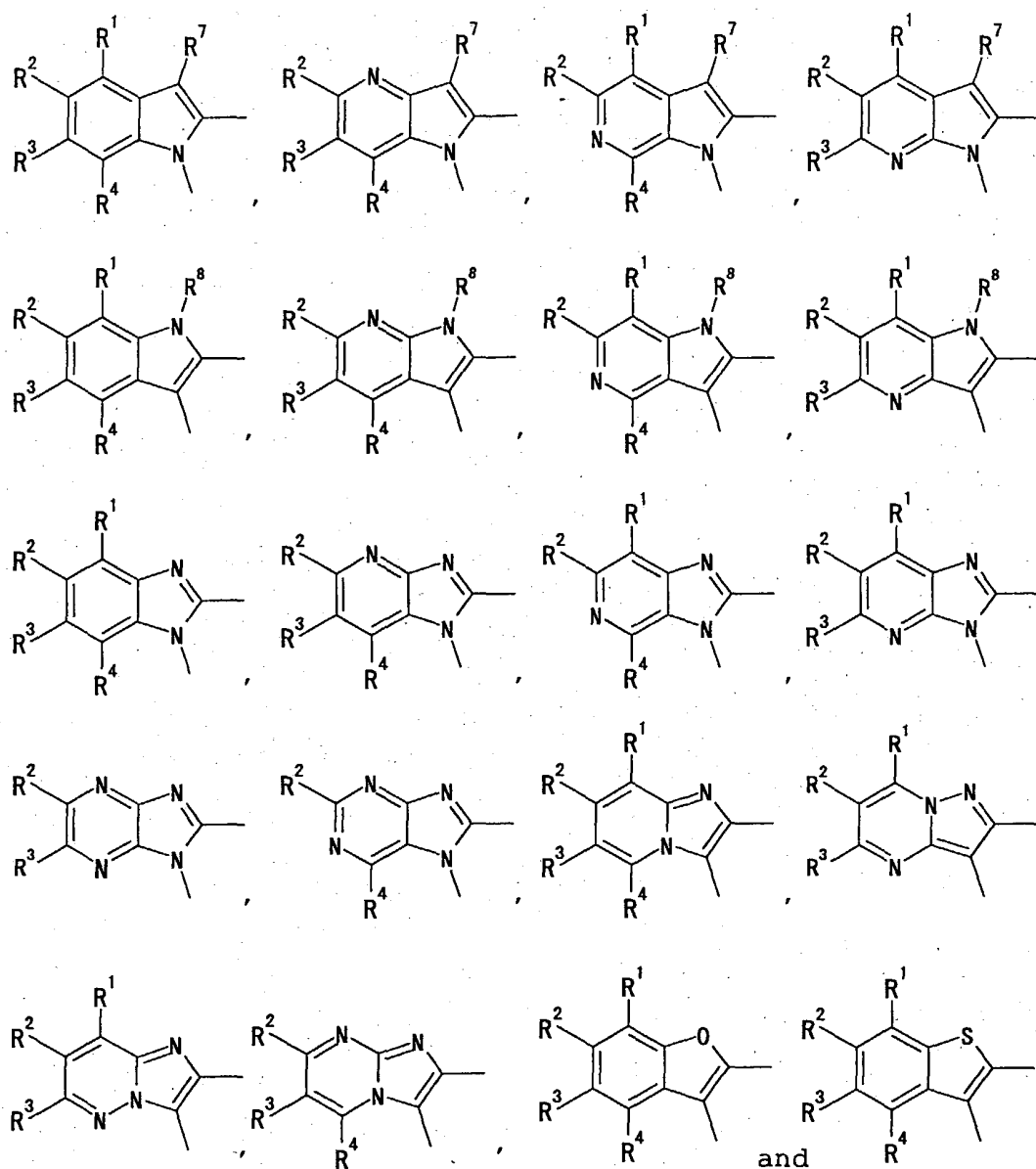
(31) The therapeutic agent of (30) above, wherein  $G^2$  is  $C(-R^2)$  and  $G^6$  is a carbon atom.

(32) The therapeutic agent of (30) or (31) above, wherein  $G^5$  is a nitrogen atom.

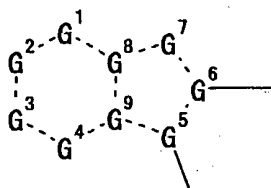
(33) The therapeutic agent of (29) above, wherein, in formula [I], the moiety



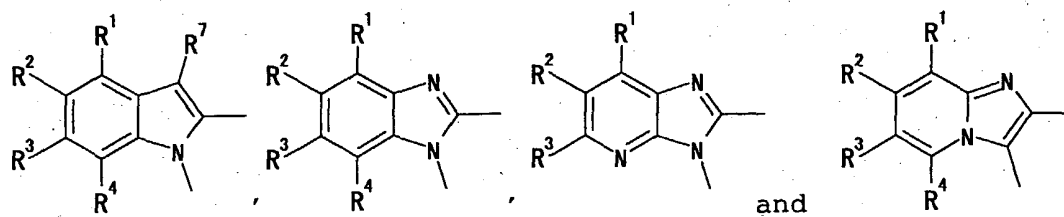
is a fused ring selected from



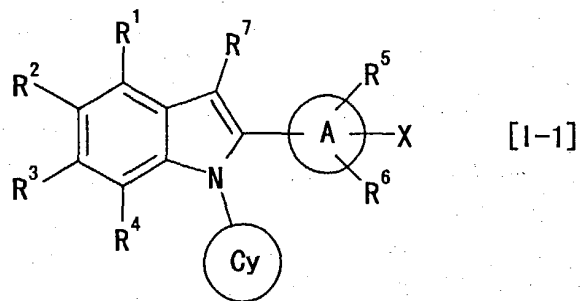
(34) The therapeutic agent of (33) above, wherein, in formula [I], the moiety



5 is a fused ring selected from

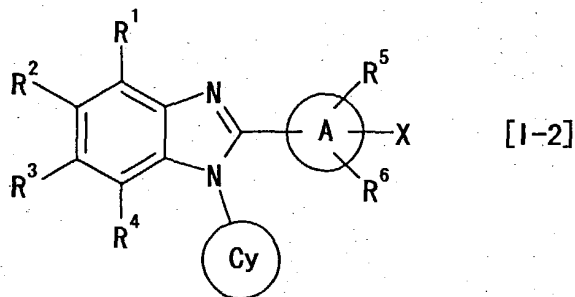


(35) The therapeutic agent of (34) above, which comprises a fused ring compound of the following formula [I-1]



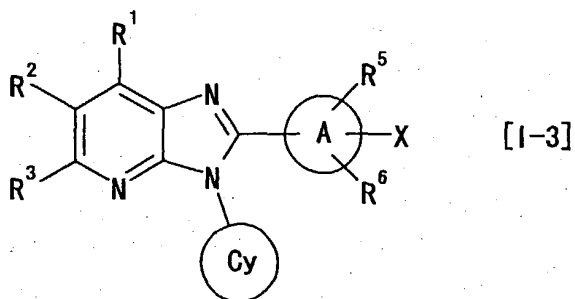
wherein each symbol is as defined in (29),  
5 or a pharmaceutically acceptable salt thereof as an active ingredient.

(36) The therapeutic agent of (34) above, which comprises a fused ring compound of the following formula [I-2]



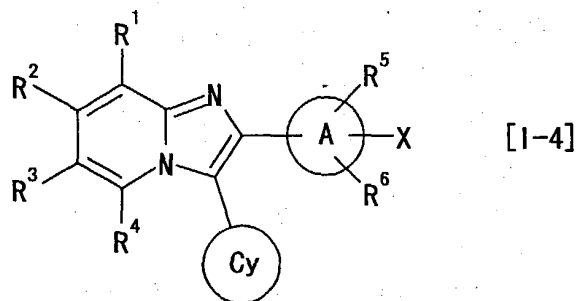
10 wherein each symbol is as defined in (29),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

(37) The therapeutic agent of (34) above, which comprises a fused ring compound of the following formula [I-3]



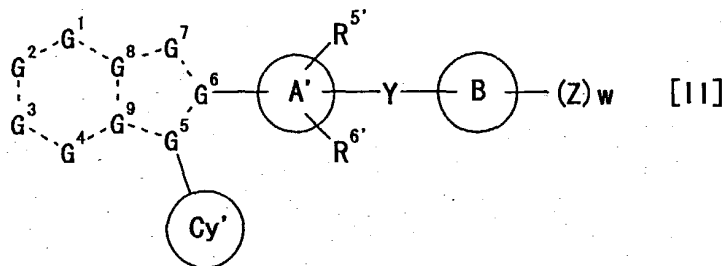
15 wherein each symbol is as defined in (29),  
or a pharmaceutically acceptable salt thereof as an active ingredient.

(38) The therapeutic agent of (34) above, which comprises a fused  
20 ring compound of the following formula [I-4]

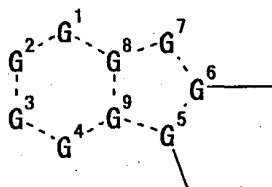


wherein each symbol is as defined in (29),  
or a pharmaceutically acceptable salt thereof as an active  
ingredient.

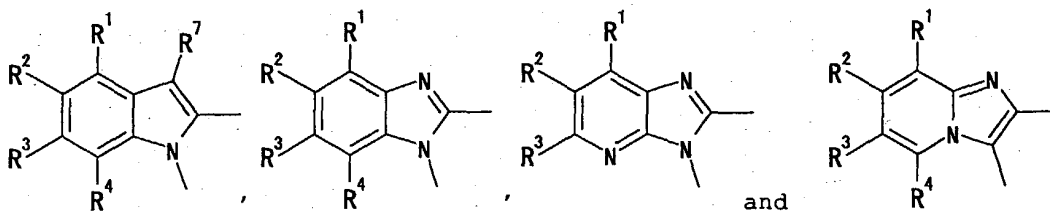
- 5 (39) The therapeutic agent of any of (29) to (38) above, wherein  
at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl,  $-\text{COOR}^{a1}$ ,  $-\text{CONR}^{a2}\text{R}^{a3}$  or  
 $-\text{SO}_2\text{R}^{a7}$  wherein  $R^{a1}$ ,  $R^{a2}$ ,  $R^{a3}$  and  $R^{a7}$  are as defined in (29).  
(40) The therapeutic agent of any of (29) to (39) above, wherein  
the ring Cy is cyclopentyl, cyclohexyl, cycloheptyl or  
10 tetrahydrothiopyranyl.  
(41) The therapeutic agent of any of (29) to (40) above, wherein  
the ring A is  $\text{C}_{6-14}$  aryl.  
(42) A fused ring compound of the following formula [II]



15 wherein  
the moiety



is a fused ring selected from





wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently,

(1) hydrogen atom,

(2)  $C_{1-6}$  alkanoyl,

(3) carboxyl,

5 (4) cyano,

(5) nitro,

(6)  $C_{1-6}$  alkyl optionally substituted by 1 to 3

substituent(s) selected from the following group A,

group A; halogen atom, hydroxyl group, carboxyl, amino,

10  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkoxy  $C_{1-6}$  alkoxy,  $C_{1-6}$   
alkoxycarbonyl and  $C_{1-6}$  alkylamino,

(7)  $-\text{COOR}^{a1}$

wherein  $R^{a1}$  is optionally substituted  $C_{1-6}$  alkyl (as  
defined above),  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally

15 substituted by 1 to 5 substituent(s) selected from the  
following group B or glucuronic acid residue,

group B; halogen atom, cyano, nitro,  $C_{1-6}$  alkyl,

halogenated  $C_{1-6}$  alkyl,  $C_{1-6}$  alkanoyl,

20  $-(\text{CH}_2)_r-\text{COOR}^{b1}$ ,  $-(\text{CH}_2)_r-\text{CONR}^{b1}\text{R}^{b2}$ ,  $-(\text{CH}_2)_r-\text{NR}^{b1}\text{R}^{b2}$ ,  
 $-(\text{CH}_2)_r-\text{NR}^{b1}-\text{COR}^{b2}$ ,  $-(\text{CH}_2)_r-\text{NHSO}_2\text{R}^{b1}$ ,  $-(\text{CH}_2)_r-\text{OR}^{b1}$ ,  
 $-(\text{CH}_2)_r-\text{SR}^{b1}$ ,  $-(\text{CH}_2)_r-\text{SO}_2\text{R}^{b1}$  and  $-(\text{CH}_2)_r-\text{SO}_2\text{NR}^{b1}\text{R}^{b2}$

wherein  $R^{b1}$  and  $R^{b2}$  are each independently  
hydrogen atom or  $C_{1-6}$  alkyl and  $r$  is 0 or an  
integer of 1 to 6,

25 (8)  $-\text{CONR}^{a2}\text{R}^{a3}$

wherein  $R^{a2}$  and  $R^{a3}$  are each independently hydrogen  
atom,  $C_{1-6}$  alkoxy or optionally substituted  $C_{1-6}$  alkyl  
(as defined above),

(9)  $-\text{C}(=\text{NR}^{a4})\text{NH}_2$

30 wherein  $R^{a4}$  is hydrogen atom or hydroxyl group,

(10)  $-\text{NHR}^{a5}$

wherein  $R^{a5}$  is hydrogen atom,  $C_{1-6}$  alkanoyl or  $C_{1-6}$   
alkylsulfonyl,

(11)  $-\text{OR}^{a6}$

35 wherein  $R^{a6}$  is hydrogen atom or optionally substituted  
 $C_{1-6}$  alkyl (as defined above),

(12)  $-\text{SO}_2\text{R}^{a7}$

wherein  $R^{a7}$  is hydroxyl group, amino,  $C_{1-6}$  alkyl or  $C_{1-6}$

alkylamino,  
(13)  $-P(=O)(OR^{a31})_2$

wherein  $R^{a31}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

or

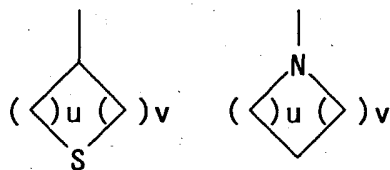
(14) heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom, and

$R^7$  is hydrogen atom or optionally substituted  $C_{1-6}$  alkyl (as defined above),

ring  $Cy'$  is

(1)  $C_{3-8}$  cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the following group C, group C; hydroxyl group, halogen atom,  $C_{1-6}$  alkyl and  $C_{1-6}$  alkoxy, or

(2)



wherein  $u$  and  $v$  are each independently an integer of 1 to 3,

ring  $A'$  is a group selected from a group consisting of phenyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, cyclohexyl, cyclohexenyl, furyl and thienyl,

$R^{5'}$  and  $R^{6'}$  are each independently

(1) hydrogen atom,  
(2) halogen atom,  
(3) optionally substituted  $C_{1-6}$  alkyl (as defined above)  
or

(4) hydroxyl group

ring B is

(1)  $C_{6-14}$  aryl,  
(2)  $C_{3-8}$  cycloalkyl or  
(3) heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a

- sulfur atom,
- each Z is independently
- (1) a group selected from the following group D,
  - (2) C<sub>6-14</sub> aryl optionally substituted by 1 to 5  
5 substituent(s) selected from the following group D,
  - (3) C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5  
substituent(s) selected from the following group D,
  - (4) C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5  
substituent(s) selected from the following group D,
  - 10 (5) heterocyclic group optionally substituted by 1 to 5  
substituent(s) selected from the following group D  
wherein the heterocyclic group has 1 to 4  
heteroatom(s) selected from an oxygen atom, a nitrogen  
atom and a sulfur atom, or
  - 15 (6) heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to  
5 substituent(s) selected from the following group D  
wherein the heterocycle C<sub>1-6</sub> alkyl is C<sub>1-6</sub> alkyl  
substituted by heterocyclic group optionally  
substituted by 1 to 5 substituent(s) selected from the  
20 group D, as defined above,  
group D:
    - (a) hydrogen atom,
    - (b) halogen atom,
    - (c) cyano,
    - 25 (d) nitro,
    - (e) optionally substituted C<sub>1-6</sub> alkyl (as defined  
above),
    - (f) -(CH<sub>2</sub>)<sub>t</sub>-COR<sup>a18</sup>,  
(hereinafter each t means independently 0 or an  
30 integer of 1 to 6),  
wherein R<sup>a18</sup> is
      - (1') optionally substituted C<sub>1-6</sub> alkyl (as  
defined above),
      - (2') C<sub>6-14</sub> aryl optionally substituted by 1 to  
35 5 substituent(s) selected from the above  
group B or
      - (3') heterocyclic group optionally  
substituted by 1 to 5 substituent(s)

selected from the above group B  
wherein the heterocyclic group has 1 to  
4 heteroatom(s) selected from an oxygen  
atom, a nitrogen atom and a sulfur atom,

5 (g)  $-(CH_2)_t-COOR^{a19}$

wherein  $R^{a19}$  is hydrogen atom, optionally  
substituted  $C_{1-6}$  alkyl (as defined above) or  
 $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1  
to 5 substituent(s) selected from the above  
10 group B,

(h)  $-(CH_2)_t-CONR^{a27}R^{a28}$

wherein  $R^{a27}$  and  $R^{a28}$  are each independently,

(1") hydrogen atom,

(2") optionally substituted  $C_{1-6}$  alkyl (as  
15 defined above),

(3")  $C_{6-14}$  aryl optionally substituted by 1 to 5  
substituent(s) selected from the above group  
B,

(4")  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted  
20 by 1 to 5 substituent(s) selected from the  
above group B,

(5") heterocyclic group optionally substituted by  
1 to 5 substituent(s) selected from the above  
group B,

25 (6") heterocycle  $C_{1-6}$  alkyl optionally  
substituted by 1 to 5 substituent(s) selected  
from the above group B,

wherein the heterocycle  $C_{1-6}$  alkyl is  $C_{1-6}$  alkyl  
substituted by heterocyclic group optionally  
30 substituted by 1 to 5 substituent(s) selected  
from the above group B, as defined above,

(7")  $C_{3-8}$  cycloalkyl optionally substituted by 1  
to 5 substituent(s) selected from the above  
group B,

35 (8")  $C_{3-8}$  cycloalkyl  $C_{1-6}$  alkyl optionally  
substituted by 1 to 5 substituent(s) selected  
from the above group B,

(9") hydroxyl group or

(10") C<sub>1-6</sub> alkoxy,

(i)  $-(CH_2)_t-C(=NR^{a33})NH_2$

wherein R<sup>a33</sup> is hydrogen atom, C<sub>1-6</sub> alkyl, hydroxyl group or C<sub>1-6</sub> alkoxy,

(j)  $-(CH_2)_t-OR^{a20}$

wherein R<sup>a20</sup> is

(1') hydrogen atom,

(2') optionally substituted C<sub>1-6</sub> alkyl (as defined above),

(3') optionally substituted C<sub>2-6</sub> alkenyl (as defined above),

(4') C<sub>2-6</sub> alkynyl optionally substituted by 1 to 3 substituent(s) selected from the above group A,

(5') C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(6') C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(7') heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(8') heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(9') C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B, or

(10') C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(k)  $-(CH_2)_t-O-(CH_2)_p-COR^{a21}$

wherein R<sup>a21</sup> is amino, C<sub>1-6</sub> alkylamino or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

and p is 0 or an integer of 1 to 6,

(l)  $-(CH_2)_t-NR^{a22}R^{a23}$

wherein  $R^{a22}$  and  $R^{a23}$  are each independently

(1') hydrogen atom,

(2') optionally substituted  $C_{1-6}$  alkyl (as  
defined above),

(3')  $C_{6-14}$  aryl optionally substituted by 1 to  
5 substituent(s) selected from the  
above group B,

(4')  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

(5') heterocycle  $C_{1-6}$  alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B or

(6') heterocyclic group optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

(m)  $-(CH_2)_t-NR^{a29}CO-R^{a24}$

wherein  $R^{a29}$  is hydrogen atom,  $C_{1-6}$  alkyl or  $C_{1-6}$   
alkanoyl, and

$R^{a24}$  is

(1') amino,

(2')  $C_{1-6}$  alkylamino,

(3') optionally substituted  $C_{1-6}$   
alkyl (as defined above),

(4')  $C_{6-14}$  aryl optionally substituted by 1  
to 5 substituent(s) selected from the  
above group B,

(5') heterocyclic group optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B, or

(6') heterocycle  $C_{1-6}$  alkyl optionally  
substituted by 1 to 5 substituent(s)  
selected from the above group B,

(n)  $-(CH_2)_t-NR^{a29}SO_2-R^{a25}$

wherein  $R^{a29}$  is as defined above, and

$R^{a25}$  is hydrogen atom, optionally  
substituted  $C_{1-6}$  alkyl (as defined above),

C<sub>6-14</sub> aryl optionally substituted by 1 to 5  
 substituent(s) selected from the above group  
 B or heterocyclic group optionally  
 substituted by 1 to 5 substituent(s) selected  
 from the above group B,  
 5  
 (o)  $-(CH_2)_t-S(O)_q-R^{a25}$   
 wherein R<sup>a25</sup> is as defined above, and q is 0,  
 1 or 2,  
 (p)  $-(CH_2)_t-SO_2-NHR^{a26}$   
 10 wherein R<sup>a26</sup> is hydrogen atom, optionally  
 substituted C<sub>1-6</sub> alkyl (as defined above),  
 C<sub>6-14</sub> aryl optionally substituted by 1 to 5  
 substituent(s) selected from the above group  
 B  
 15 or heterocyclic group optionally substituted  
 by 1 to 5 substituent(s) selected from the  
 above group B,  
 and  
 (q) heterocyclic group having 1 to 4  
 20 heteroatom(s) selected from an oxygen atom,  
 a nitrogen atom and a sulfur atom,  
 w is an integer of 1 to 3, and  
 Y is  
 (1) a single bond,  
 25 (2) C<sub>1-6</sub> alkylene,  
 (3) C<sub>2-6</sub> alkenylene,  
 (4)  $-(CH_2)_m-O-(CH_2)_n-$ ,  
 (hereinafter m and n are each independently 0  
 or an integer of 1 to 6),  
 30 (5)  $-CO-$ ,  
 (6)  $-CO_2-(CH_2)_n-$ ,  
 (7)  $-CONH-(CH_2)_n-NH-$ ,  
 (8)  $-NHCO_2-$ ,  
 (9)  $-NHCONH-$ ,  
 35 (10)  $-O-(CH_2)_n-CO-$ ,  
 (11)  $-O-(CH_2)_n-O-$ ,  
 (12)  $-SO_2-$ ,  
 (13)  $-(CH_2)_m-NR^{a12}-(CH_2)_n-$

wherein  $R^{a12}$  is

- (1') hydrogen atom,
- (2') optionally substituted  $C_{1-6}$  alkyl (as defined above),
- (3')  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (4')  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (5')  $-COR^{b5}$

wherein  $R^{b5}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

- (6')  $-COOR^{b5}$  ( $R^{b5}$  is as defined above) or
- (7')  $-SO_2R^{b5}$  ( $R^{b5}$  is as defined above),
- (14)  $-NR^{a12}CO-$  ( $R^{a12}$  is as defined above),
- (15)  $-CONR^{a13}-(CH_2)_n-$

wherein  $R^{a13}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

- (16)  $-CONH-CHR^{a14}-$

wherein  $R^{a14}$  is  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

- (17)  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$

wherein  $R^{a15}$  and  $R^{a16}$  are each independently

- (1') hydrogen atom,
- (2') carboxyl,
- (3')  $C_{1-6}$  alkyl,
- (4')  $-OR^{b6}$



wherein  $R^{b6}$  is  $C_{1-6}$  alkyl or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl, or

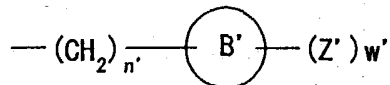
(5')  $-NHR^{b7}$

wherein  $R^{b7}$  is hydrogen atom,  $C_{1-6}$  alkyl,  $C_{1-6}$  alkanoyl or  $C_{6-14}$  aryl  $C_{1-6}$  alkyloxycarbonyl,

or

$R^{a15}$  is optionally

(6')



wherein  $n'$ , ring  $B'$ ,  $Z'$  and  $w'$  are the same as the above-mentioned  $n$ , ring  $B$ ,  $Z$  and  $w$ , respectively, and may be the same as or different from the respective counterparts,

(18)  $-(CH_2)_n-NR^{a12}-CHR^{a15}-$  ( $R^{a12}$  and  $R^{a15}$  are each as defined above),

(19)  $-NR^{a17}SO_2-$

wherein  $R^{a17}$  is hydrogen atom or  $C_{1-6}$  alkyl,

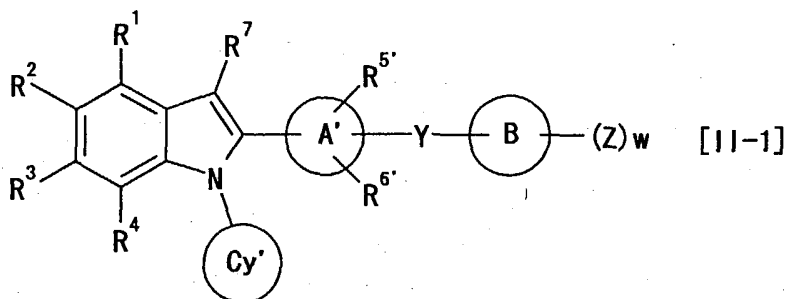
(20)  $-S(O)_e-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  ( $e$  is 0, 1 or 2,  $R^{a15}$  and  $R^{a16}$  are each as defined above),

or

(21)  $-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  ( $R^{a15}$  and  $R^{a16}$  are each as defined above),

or a pharmaceutically acceptable salt thereof.

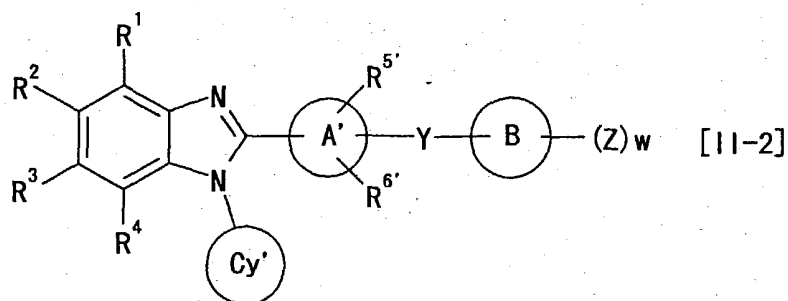
(43) The fused ring compound of (42) above, which is represented by the following formula [II-1]



wherein each symbol is as defined in (42),

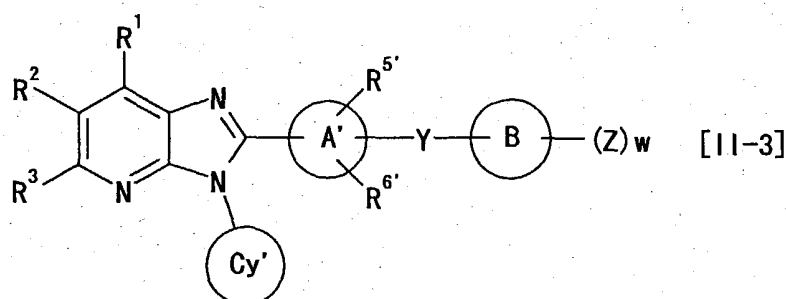
or a pharmaceutically acceptable salt thereof.

(44) The fused ring compound of (42) above, which is represented by the following formula [II-2]



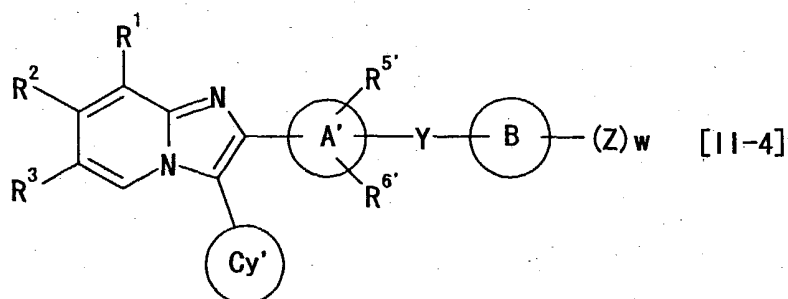
wherein each symbol is as defined in (42),  
or a pharmaceutically acceptable salt thereof.

(45) The fused ring compound of (42) above, which is represented  
5 by the following formula [II-3]



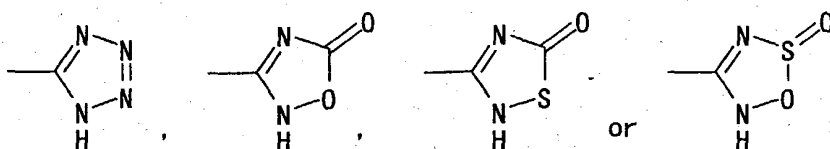
wherein each symbol is as defined in (42),  
or a pharmaceutically acceptable salt thereof.

(46) The fused ring compound of (42) above, which is represented  
10 by the following formula [II-4]



wherein each symbol is as defined in (42),  
or a pharmaceutically acceptable salt thereof.

(47) The fused ring compound of any of (42) to (46) above,  
15 wherein at least one of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> is carboxyl, -COOR<sup>a1</sup>,  
-CONR<sup>a2</sup>R<sup>a3</sup>, -SO<sub>2</sub>R<sup>a7</sup> (wherein R<sup>a1</sup>, R<sup>a2</sup>, R<sup>a3</sup> and R<sup>a7</sup> are as defined in  
(42)),



or a pharmaceutically acceptable salt thereof.

(48) The fused ring compound of (47) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl,  $-\text{COOR}^{a1}$  or  $-\text{SO}_2\text{R}^{a7}$  wherein  $R^{a1}$  and  $R^{a7}$  are as defined in (42), or a pharmaceutically acceptable salt thereof.

(49) The fused ring compound of (48) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl or  $-\text{COOR}^{a1}$  wherein  $R^{a1}$  is as defined in (42), or a pharmaceutically acceptable salt thereof.

(50) The fused ring compound of (49) above, wherein  $R^2$  is carboxyl and  $R^1$ ,  $R^3$  and  $R^4$  are hydrogen atoms, or a pharmaceutically acceptable salt thereof.

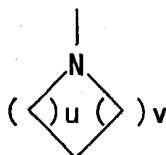
(51) The fused ring compound of any of (42) to (46) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is carboxyl or  $-\text{COOR}^{a1}$  wherein  $R^{a1}$  is glucuronic acid residue, or a pharmaceutically acceptable salt thereof.

(52) The fused ring compound of any of (42) to (46) above, wherein at least one of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom, or a pharmaceutically acceptable salt thereof.

(53) The fused ring compound of any of (42) to (52) above, wherein the ring  $\text{Cy}'$  is cyclopentyl, cyclohexyl, cycloheptyl or tetrahydrothiopyranyl, or a pharmaceutically acceptable salt thereof.

(54) The fused ring compound of (42) above, wherein the ring  $\text{Cy}'$  is cyclopentyl, cyclohexyl or cycloheptyl, or a pharmaceutically acceptable salt thereof.

(55) The fused ring compound of any of (42) to (52) above, wherein the ring  $\text{Cy}'$  is



wherein each symbol is as defined in (42), or a pharmaceutically acceptable salt thereof.

(56) The fused ring compound of any of (42) to (55) above, wherein the ring A' is phenyl, pyridyl, pyrazinyl, pyrimidinyl or  
5 pyridazinyl, or a pharmaceutically acceptable salt thereof.

(57) The fused ring compound of (56) above, wherein the ring A' is phenyl or pyridyl, or a pharmaceutically acceptable salt thereof.

(58) The fused ring compound of (57) above, wherein the ring A'  
10 is phenyl, or a pharmaceutically acceptable salt thereof.

(59) The fused ring compound of any of (42) to (58) above, wherein at least one substituent optionally substituted by group A is a substituent substituted by C<sub>1-6</sub> alkoxy C<sub>1-6</sub> alkoxy, or a pharmaceutically acceptable salt thereof.

15 (60) The fused ring compound of any of (42) to (59) above, wherein the Y is  $-(CH_2)_m-O-(CH_2)_n-$ ,  $-NHCO_2-$ ,  $-CONH-CHR^{a14}-$ ,  $-(CH_2)_m-NR^{a12}-(CH_2)_n-$ ,  $-CONR^{a13}-(CH_2)_n-$ ,  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  or  $-(CH_2)_n-NR^{a12}-CHR^{a15}-$  (wherein each symbol is as defined in (42)), or a pharmaceutically acceptable salt thereof.

20 (61) The fused ring compound of (42) above, wherein the Y is  $-(CH_2)_m-O-(CH_2)_n-$  or  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  (wherein each symbol is as defined in (42)), or a pharmaceutically acceptable salt thereof.

(62) The fused ring compound of (61) above, wherein the Y is  
25  $-(CH_2)_m-O-(CH_2)_n-$  wherein each symbol is as defined in (42), or a pharmaceutically acceptable salt thereof.

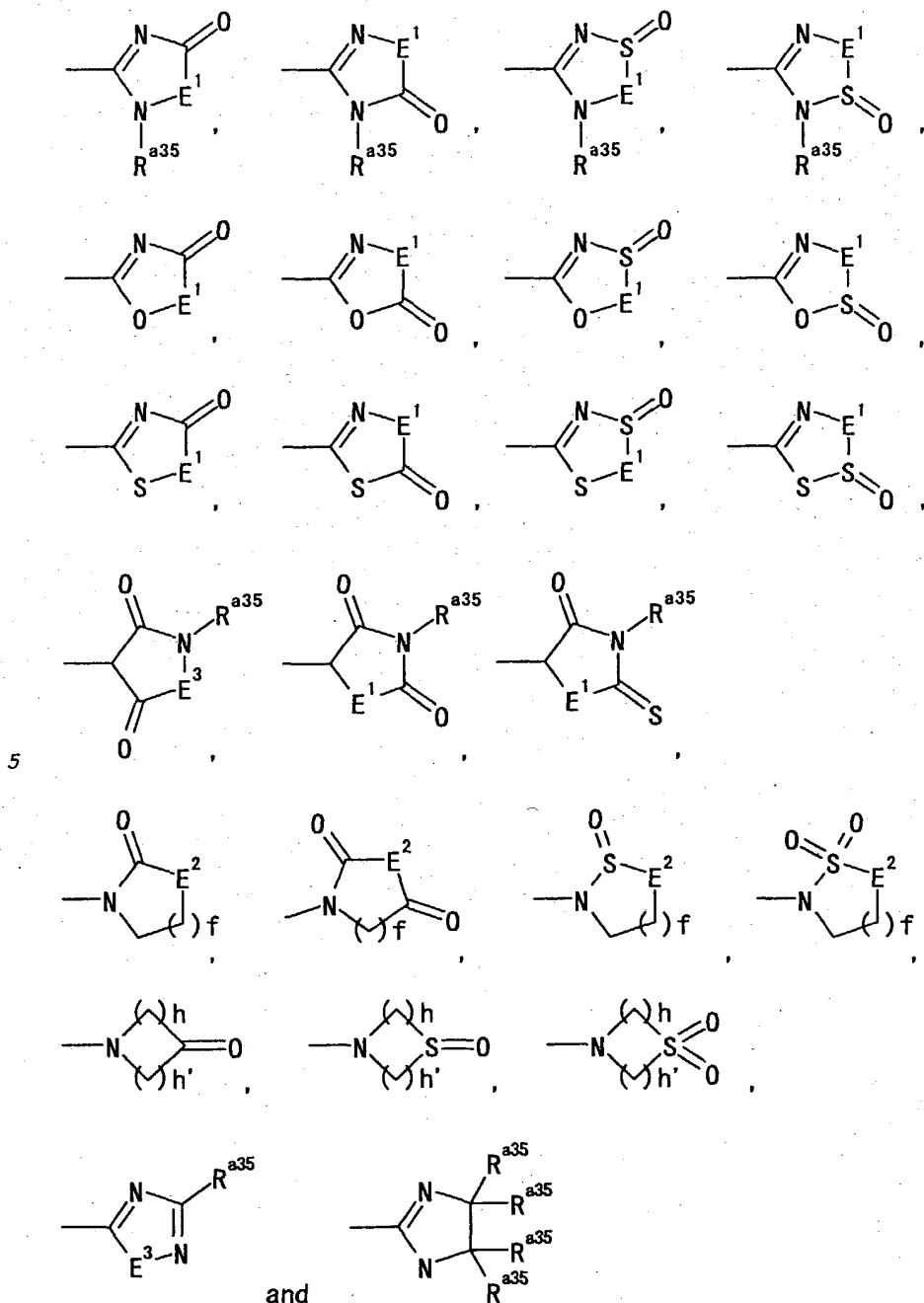
(63) The fused ring compound of any of (42) to (59) above, wherein the Y is  $-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  (wherein each symbol is as defined in (42)), or a pharmaceutically acceptable salt thereof.

30 (64) The fused ring compound of any of (42) to (63) above, wherein the R<sup>2</sup> is carboxyl, R<sup>1</sup>, R<sup>3</sup> and R<sup>4</sup> are hydrogen atoms, the ring Cy' is cyclopentyl, cyclohexyl or cycloheptyl, and the ring A' is phenyl, or a pharmaceutically acceptable salt thereof.

(65) The fused ring compound of any of (42) to (64) above,  
35 wherein at least one group represented by Z is heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the group D, or a pharmaceutically acceptable salt thereof.

(66) The fused ring compound of any of (42) to (64) above,

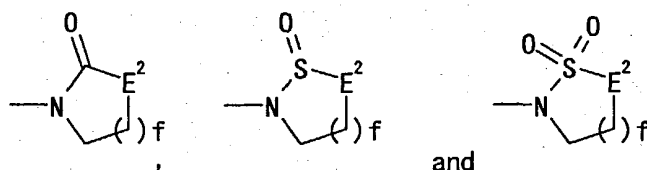
wherein at least one group represented by Z is heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the group D, wherein said heterocyclic group is selected from the following groups:



10 wherein  $E^1$  is an oxygen atom, a sulfur atom or  $N(-R^{a35})$ ,  $E^2$  is an oxygen atom,  $CH_2$  or  $N(-R^{a35})$ ,  $E^3$  is an oxygen atom or a sulfur atom, wherein each  $R^{a35}$  is independently hydrogen atom or  $C_{1-6}$  alkyl,  $f$  is an integer of 1 to 3, and  $h$  and  $h'$  are the same or different

and each is an integer of 1 to 3, or a pharmaceutically acceptable salt thereof.

- (67) The fused ring compound of (66) above, wherein at least one group represented by Z is heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the group D, wherein said heterocyclic group is selected from the following groups:



wherein each symbol is as defined in (66), or a pharmaceutically acceptable salt thereof.

- (68) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is  $-(CH_2)_t-CONR^{a27}R^{a28}$  wherein each symbol is as defined in (42), and at least one of  $R^{a27}$  and  $R^{a28}$  is  $C_{1-6}$  alkoxy, or a pharmaceutically acceptable salt thereof.

- (69) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is  $-(CH_2)_t-C(=NR^{a33})NH_2$  wherein each symbol is as defined in (42), and  $R^{a33}$  is hydroxyl group or  $C_{1-6}$  alkoxy, or a pharmaceutically acceptable salt thereof.

- (70) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is  $-(CH_2)_t-O-(CH_2)_p-COR^{a21}$  wherein each symbol is as defined in (42), and  $R^{a21}$  is amino, or a pharmaceutically acceptable salt thereof.

- (71) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is  $-(CH_2)_t-NR^{a29}CO-R^{a24}$  wherein each symbol is as defined in (42), and  $R^{a24}$  is amino or

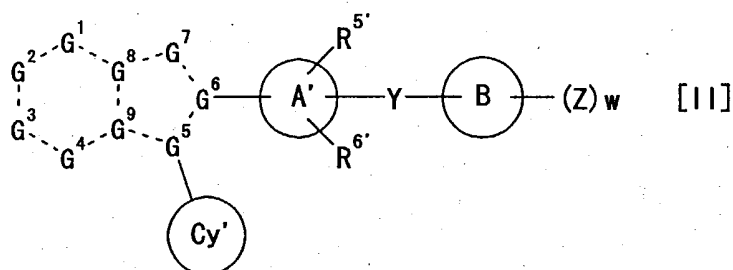
$C_{1-6}$  alkylamino, or a pharmaceutically acceptable salt thereof.

- (72) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is  $-(CH_2)_t-NR^{a22}R^{a23}$  wherein each symbol is as defined in (42), and at least one of  $R^{a22}$  and  $R^{a23}$  is heterocyclic group optionally substituted

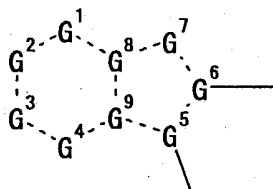
by 1 to 5 substituent(s) selected from the group B, or a pharmaceutically acceptable salt thereof.

(73) The fused ring compound of any of (42) to (64) above, wherein at least one group represented by group D is heterocyclic group having 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom, or a pharmaceutically acceptable salt thereof.

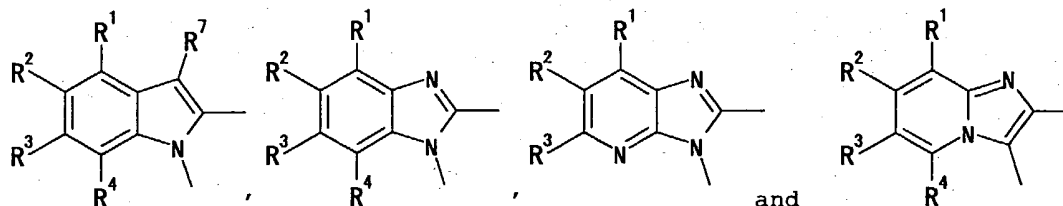
(74) The fused ring compound of (42) above, which is represented by the following formula [II]



wherein  
the moiety



is a fused ring selected from



wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently,

- (1) hydrogen atom,
- (2)  $C_{1-6}$  alkanoyl,
- (3) carboxyl,
- (4) cyano,
- (5) nitro,

(6)  $C_{1-6}$  alkyl optionally substituted by 1 to 3 substituent(s) selected from the following group A, group A; halogen atom, hydroxyl group, carboxyl, amino,

$C_{1-6}$  alkoxy,  $C_{1-6}$  alkoxy carbonyl and  $C_{1-6}$

alkylamino,

(7)  $-\text{COOR}^{\text{a1}}$

wherein  $\text{R}^{\text{a1}}$  is optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above) or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the following group B,

group B; halogen atom, cyano, nitro,  $\text{C}_{1-6}$  alkyl,

halogenated  $\text{C}_{1-6}$  alkyl,  $\text{C}_{1-6}$  alkanoyl,

$-(\text{CH}_2)_r-\text{COOR}^{\text{b1}}$ ,  $-(\text{CH}_2)_r-\text{CONR}^{\text{b1}}\text{R}^{\text{b2}}$ ,  $-(\text{CH}_2)_r-\text{NR}^{\text{b1}}\text{R}^{\text{b2}}$ ,

$-(\text{CH}_2)_r-\text{NR}^{\text{b1}}-\text{COR}^{\text{b2}}$ ,  $-(\text{CH}_2)_r-\text{NHSO}_2\text{R}^{\text{b1}}$ ,  $-(\text{CH}_2)_r-\text{OR}^{\text{b1}}$ ,

$-(\text{CH}_2)_r-\text{SR}^{\text{b1}}$ ,  $-(\text{CH}_2)_r-\text{SO}_2\text{R}^{\text{b1}}$  and  $-(\text{CH}_2)_r-\text{SO}_2\text{NR}^{\text{b1}}\text{R}^{\text{b2}}$

wherein  $\text{R}^{\text{b1}}$  and  $\text{R}^{\text{b2}}$  are each independently hydrogen atom or  $\text{C}_{1-6}$  alkyl and  $r$  is 0 or an integer of 1 to 6,

(8)  $-\text{CONR}^{\text{a2}}\text{R}^{\text{a3}}$

wherein  $\text{R}^{\text{a2}}$  and  $\text{R}^{\text{a3}}$  are each independently hydrogen atom,  $\text{C}_{1-6}$  alkoxy or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

(9)  $-\text{C}(=\text{NR}^{\text{a4}})\text{NH}_2$

wherein  $\text{R}^{\text{a4}}$  is hydrogen atom or hydroxyl group,

(10)  $-\text{NHR}^{\text{a5}}$

wherein  $\text{R}^{\text{a5}}$  is hydrogen atom,  $\text{C}_{1-6}$  alkanoyl or  $\text{C}_{1-6}$  alkylsulfonyl,

(11)  $-\text{OR}^{\text{a6}}$

wherein  $\text{R}^{\text{a6}}$  is hydrogen atom or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

(12)  $-\text{SO}_2\text{R}^{\text{a7}}$

wherein  $\text{R}^{\text{a7}}$  is hydroxyl group, amino,  $\text{C}_{1-6}$  alkyl or  $\text{C}_{1-6}$  alkylamino

or

(13)  $-\text{P}(=\text{O})(\text{OR}^{\text{a31}})_2$

wherein  $\text{R}^{\text{a31}}$  is hydrogen atom, optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above) or  $\text{C}_{6-14}$  aryl  $\text{C}_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B, and

$\text{R}^7$  is hydrogen atom or optionally substituted  $\text{C}_{1-6}$  alkyl (as defined above),

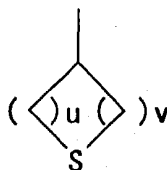
ring Cy' is



- (1) C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5  
substituent(s) selected from the following group C,  
group C; hydroxyl group, halogen atom, C<sub>1-6</sub> alkyl and  
C<sub>1-6</sub> alkoxy, or

5

(2)



wherein u and v are each independently an integer  
of 1 to 3,

ring A' is a group selected from a group consisting of  
10 phenyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl,  
cyclohexyl, cyclohexenyl, furyl and thienyl,

R<sup>5'</sup> and R<sup>6'</sup> are each independently

- (1) hydrogen atom,  
(2) halogen atom,  
15 (3) optionally substituted C<sub>1-6</sub> alkyl (as defined above)  
or  
(4) hydroxyl group

ring B is

- (1) C<sub>6-14</sub> aryl,  
20 (2) C<sub>3-8</sub> cycloalkyl or  
(3) heterocyclic group having 1 to 4 heteroatom(s)

selected from an oxygen atom, a nitrogen atom and a sulfur atom,  
each Z is independently

- (1) a group selected from the following group D,  
25 (2) C<sub>6-14</sub> aryl optionally substituted by 1 to 5  
substituent(s) selected from the following group D,  
(3) C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5  
substituent(s) selected from the following group D,  
(4) C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5  
30 substituent(s) selected from the following group D or  
(5) heterocyclic group optionally substituted by 1 to 5  
substituent(s) selected from the following group D  
wherein the heterocyclic group has 1 to 4  
heteroatom(s) selected from an oxygen atom, a nitrogen  
35 atom and a sulfur atom,

group D:

- (a) hydrogen atom,
- (b) halogen atom,
- (c) cyano,
- (d) nitro,
- (e) optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- (f)  $-(CH_2)_t-COR^{a18}$ ,

(hereinafter each t means independently 0 or an integer of 1 to 6),

wherein R<sup>a18</sup> is

- (1') optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- (2') C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or

- (3') heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B wherein the heterocyclic group has 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom,

- (g)  $-(CH_2)_t-COOR^{a19}$

wherein R<sup>a19</sup> is hydrogen atom, optionally substituted C<sub>1-6</sub> alkyl (as defined above) or C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

- (h)  $-(CH_2)_t-CONR^{a27}R^{a28}$

wherein R<sup>a27</sup> and R<sup>a28</sup> are each independently,

- (1'') hydrogen atom,
- (2'') optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- (3'') C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (4'') C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the

- above group B,
- (5") heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 5 (6") heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- wherein the heterocycle C<sub>1-6</sub> alkyl is C<sub>1-6</sub> alkyl substituted by heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B, as defined above,
- 10 (7") C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B, or
- 15 (8") C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (i)  $-(CH_2)_t-C(=NR^{a33})NH_2$   
wherein R<sup>a33</sup> is hydrogen atom or C<sub>1-6</sub> alkyl,
- 20 (j)  $-(CH_2)_t-OR^{a20}$   
wherein R<sup>a20</sup> is
- (1') hydrogen atom,
- (2') optionally substituted C<sub>1-6</sub> alkyl (as defined above),
- 25 (3') optionally substituted C<sub>2-6</sub> alkenyl (as defined above),
- (4') C<sub>2-6</sub> alkynyl optionally substituted by 1 to 3 substituent(s) selected from the above group A,
- 30 (5') C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (6') C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- 35 (7') heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

- (8') heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (9') C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B, or
- (10') C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (k)  $-(CH_2)_t-O-(CH_2)_p-COR^{a21}$   
wherein R<sup>a21</sup> is C<sub>1-6</sub> alkylamino or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,  
and p is 0 or an integer of 1 to 6,
- (l)  $-(CH_2)_t-NR^{a22}R^{a23}$   
wherein R<sup>a22</sup> and R<sup>a23</sup> are each independently  
(1') hydrogen atom,  
(2') optionally substituted C<sub>1-6</sub> alkyl (as defined above),  
(3') C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,  
(4') C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B or  
(5') heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,
- (m)  $-(CH_2)_t-NR^{a29}CO-R^{a24}$   
wherein R<sup>a29</sup> is hydrogen atom, C<sub>1-6</sub> alkyl or C<sub>1-6</sub> alkanoyl, R<sup>a24</sup> is optionally substituted C<sub>1-6</sub> alkyl (as defined above), C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(n)  $-(CH_2)_t-NHSO_2-R^{a25}$

wherein  $R^{a25}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group

B

or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(o)  $-(CH_2)_t-S(O)_q-R^{a25}$

wherein  $R^{a25}$  is as defined above, and q is 0, 1 or 2,

and

(p)  $-(CH_2)_t-SO_2-NHR^{a26}$

wherein  $R^{a26}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from the above group B,

is an integer of 1 to 3, and

is

(1) a single bond,

(2)  $C_{1-6}$  alkylene,

(3)  $C_{2-6}$  alkenylene,

(4)  $-(CH_2)_m-O-(CH_2)_n-$ ,

(hereinafter m and n are each independently 0 or an integer of 1 to 6),

(5)  $-CO-$ ,

(6)  $-CO_2-(CH_2)_n-$ ,

(7)  $-CONH-(CH_2)_n-NH-$ ,

(8)  $-NHCO_2-$ ,

(9)  $-NHCONH-$ ,

(10)  $-O-(CH_2)_n-CO-$ ,

(11)  $-O-(CH_2)_n-O-$ ,

(12)  $-SO_2-$ ,

(13)  $-(CH_2)_m-NR^{a12}-(CH_2)_n-$

wherein  $R^{a12}$  is

(1') hydrogen atom,

(2') optionally substituted  $C_{1-6}$  alkyl (as defined above),

(3')  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(4')  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(5')  $-COR^{b5}$

wherein  $R^{b5}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above),  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(6')  $-COOR^{b5}$  ( $R^{b5}$  is as defined above) or

(7')  $-SO_2R^{b5}$  ( $R^{b5}$  is as defined above),

(14)  $-NR^{a12}CO-$  ( $R^{a12}$  is as defined above),

(15)  $-CONR^{a13}-(CH_2)_n-$

wherein  $R^{a13}$  is hydrogen atom, optionally substituted  $C_{1-6}$  alkyl (as defined above) or  $C_{6-14}$  aryl  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(16)  $-CONH-CHR^{a14}-$

wherein  $R^{a14}$  is  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from the above group B,

(17)  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$

wherein  $R^{a15}$  and  $R^{a16}$  are each independently

(1') hydrogen atom,

(2') carboxyl,

(3')  $C_{1-6}$  alkyl,

(4')  $-OR^{b6}$

wherein  $R^{b6}$  is  $C_{1-6}$  alkyl or  $C_{6-14}$  aryl  $C_{1-6}$

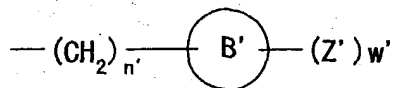
alkyl,

or

(5')  $\text{-NHR}^{b7}$

wherein  $R^{b7}$  is hydrogen atom,  $C_{1-6}$  alkyl,  $C_{1-6}$  alkanoyl or  $C_{6-14}$  aryl  $C_{1-6}$  alkyloxycarbonyl, or  $R^{a15}$  is optionally

(6')



wherein  $n'$ , ring  $B'$ ,  $Z'$  and  $w'$  are the same as the above-mentioned  $n$ , ring  $B$ ,  $Z$  and  $w$ , respectively, and may be the same as or different from the respective counterparts,

(18)  $\text{---}(\text{CH}_2)_n\text{---NR}^{a12}\text{---CHR}^{a15}\text{---}$  ( $R^{a12}$  and  $R^{a15}$  are each as defined above),

(19)  $\text{---NR}^{a17}\text{SO}_2\text{---}$

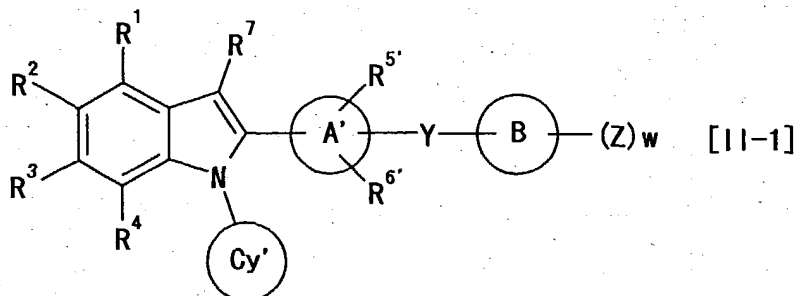
wherein  $R^{a17}$  is hydrogen atom or  $C_{1-6}$  alkyl

or

(20)  $\text{---S(O)}_e\text{---}(\text{CH}_2)_m\text{---CR}^{a15}\text{R}^{a16}\text{---}(\text{CH}_2)_n\text{---}$  ( $e$  is 0, 1 or 2,  $R^{a15}$  and  $R^{a16}$  are each as defined above),

or a pharmaceutically acceptable salt thereof.

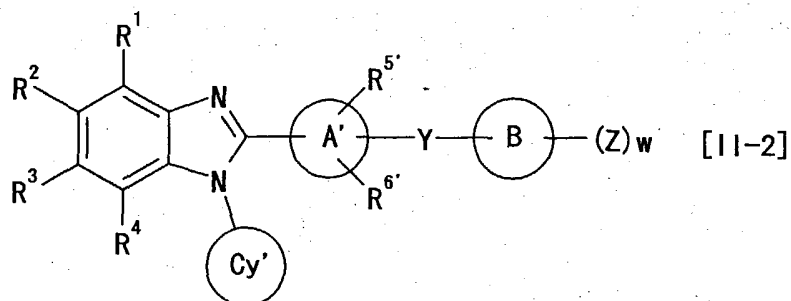
(75) The fused ring compound of (74) above, which is represented by the following formula [II-1]



wherein each symbol is as defined in (74),

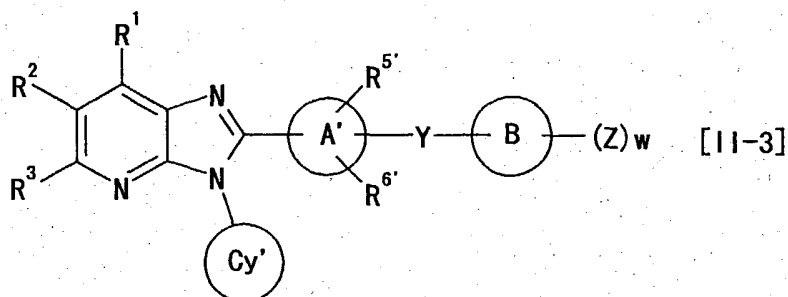
or a pharmaceutically acceptable salt thereof.

(76) The fused ring compound of (74) above, which is represented by the following formula [II-2]



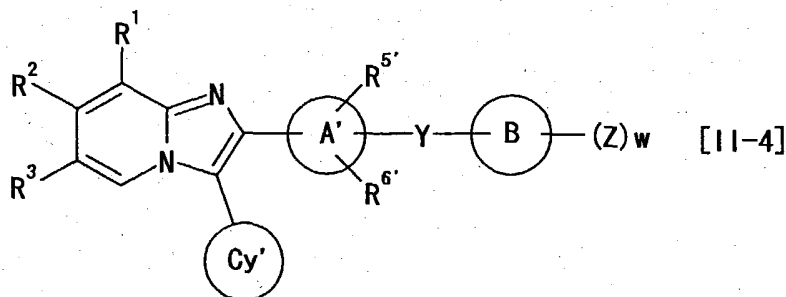
wherein each symbol is as defined in (74),  
or a pharmaceutically acceptable salt thereof.

(77) The fused ring compound of (74) above, which is represented  
5 by the following formula [II-3]



wherein each symbol is as defined in (74),  
or a pharmaceutically acceptable salt thereof.

(78) The fused ring compound of (74) above, which is represented  
10 by the following formula [II-4]



wherein each symbol is as defined in (74),  
or a pharmaceutically acceptable salt thereof.

(79) The fused ring compound of any of (74) to (78) above,  
15 wherein at least one of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> is carboxyl, -COOR<sup>a1</sup> or  
-SO<sub>2</sub>R<sup>a7</sup> wherein R<sup>a1</sup> and R<sup>a7</sup> are as defined in (74), or a  
pharmaceutically acceptable salt thereof.

(80) The fused ring compound of (79) above, wherein at least one  
of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> is carboxyl or -COOR<sup>a1</sup> wherein R<sup>a1</sup> is as  
20 defined in (74), or a pharmaceutically acceptable salt thereof.



- (81) The fused ring compound of (80) above, wherein  $R^2$  is carboxyl and  $R^1$ ,  $R^3$  and  $R^4$  are hydrogen atoms, or a pharmaceutically acceptable salt thereof.
- (82) The fused ring compound of any of (74) to (81) above,  
5 wherein the ring  $Cy'$  is cyclopentyl, cyclohexyl, cycloheptyl or tetrahydrothiopyranyl, or a pharmaceutically acceptable salt thereof.
- (83) The fused ring compound of (82) above, wherein the ring  $Cy'$  is cyclopentyl, cyclohexyl or cycloheptyl, or a pharmaceutically  
10 acceptable salt thereof.
- (84) The fused ring compound of any of (74) to (83) above, wherein the ring  $A'$  is phenyl, pyridyl, pyrazinyl, pyrimidinyl or pyridazinyl, or a pharmaceutically acceptable salt thereof.
- (85) The fused ring compound of (84) above, wherein the ring  $A'$   
15 is phenyl or pyridyl, or a pharmaceutically acceptable salt thereof.
- (86) The fused ring compound of (85) above, wherein the ring  $A'$  is phenyl, or a pharmaceutically acceptable salt thereof.
- (87) The fused ring compound of any of (74) to (86) above,  
20 wherein the  $Y$  is  $-(CH_2)_m-O-(CH_2)_n-$ ,  $-NHCO_2-$ ,  $-CONH-CHR^{a14}-$ ,  
 $-(CH_2)_m-NR^{a12}-(CH_2)_n-$ ,  $-CONR^{a13}-(CH_2)_n-$ ,  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  or  
 $-(CH_2)_n-NR^{a12}-CHR^{a15}-$  (wherein each symbol is as defined in (74)),  
or a pharmaceutically acceptable salt thereof.
- (88) The fused ring compound of (87) above, wherein the  $Y$  is  
25  $-(CH_2)_m-O-(CH_2)_n-$  or  $-O-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$  (wherein each symbol is as defined in (74)), or a pharmaceutically acceptable salt thereof.
- (89) The fused ring compound of (88) above, wherein the  $Y$  is  
 $-(CH_2)_m-O-(CH_2)_n-$  wherein each symbol is as defined in (74), or a  
30 pharmaceutically acceptable salt thereof.
- (90) The fused ring compound of any of (74) to (89) above, wherein the  $R^2$  is carboxyl,  $R^1$ ,  $R^3$  and  $R^4$  are hydrogen atoms, the ring  $Cy'$  is cyclopentyl, cyclohexyl or cycloheptyl, and the ring  $A'$  is phenyl, or a pharmaceutically acceptable salt thereof.
- 35 (91) The fused ring compound of the formula [I] or a pharmaceutically acceptable salt thereof, which is selected from the group consisting of

ethyl 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 1),  
 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 2),  
 5 ethyl 1-cyclohexyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylate (Example 3),  
 ethyl 2-[4-(2-bromo-5-chlorobenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 4),  
 ethyl 2-{4-[2-(4-chlorophenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 5),  
 10 2-{4-[2-(4-chlorophenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 6),  
 ethyl 2-[4-(2-bromo-5-methoxybenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 7),  
 15 ethyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 8),  
 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 9),  
 ethyl 1-cyclohexyl-2-{4-[(E)-2-phenylvinyl]phenyl}benzimidazole-  
 20 5-carboxylate (Example 10),  
 1-cyclohexyl-2-{4-[(E)-2-phenylvinyl]phenyl}benzimidazole-5-carboxylic acid (Example 11),  
 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 12),  
 25 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxamide (Example 13),  
 2-(4-benzyloxyphenyl)-5-cyano-1-cyclopentylbenzimidazole (Example 14),  
 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxamide  
 30 oxime (Example 15),  
 ethyl 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylate (Example 16),  
 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylic acid (Example 17),  
 35 ethyl 1-cyclohexyl-2-(2-fluoro-4-hydroxyphenyl)benzimidazole-5-carboxylate (Example 18),  
 ethyl 2-{4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 19),

2-{4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl}-1-  
 cyclohexylbenzimidazole-5-carboxylic acid (Example 20),  
 ethyl 1-cyclopentyl-2-(4-nitrophenyl)benzimidazole-5-carboxylate  
 (Example 21),  
 5 ethyl 2-(4-aminophenyl)-1-cyclopentylbenzimidazole-5-carboxylate  
 (Example 22),  
 ethyl 2-(4-benzoylamino-phenyl)-1-cyclopentylbenzimidazole-5-  
 carboxylate (Example 23),  
 2-(4-benzoylamino-phenyl)-1-cyclopentylbenzimidazole-5-carboxylic  
 10 acid (Example 24),  
 ethyl 2-{4-[3-(3-chlorophenyl)phenoxy]phenyl}-1-  
 cyclohexylbenzimidazole-5-carboxylate (Example 25),  
 2-{4-[3-(3-chlorophenyl)phenoxy]phenyl}-1-  
 cyclohexylbenzimidazole-5-carboxylic acid (Example 26),  
 15 ethyl 2-[4-(3-acetoxyphenyloxy)phenyl]-1-  
 cyclohexylbenzimidazole-5-carboxylate (Example 27),  
 ethyl 1-cyclohexyl-2-[4-(3-hydroxyphenyloxy)phenyl]-  
 benzimidazole-5-carboxylate (Example 28),  
 ethyl 1-cyclohexyl-2-{4-[3-(4-pyridylmethoxy)phenyloxy]phenyl}-  
 20 benzimidazole-5-carboxylate (Example 29),  
 1-cyclohexyl-2-{4-[3-(4-pyridylmethoxy)phenyloxy]phenyl}-  
 benzimidazole-5-carboxylic acid (Example 30),  
 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole (Example 31),  
 ethyl 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-  
 25 carboxylate (Example 32),  
 2-(4-benzyloxyphenyl)-1-cyclopentyl-N,N-dimethylbenzimidazole-5-  
 carboxamide (Example 33),  
 2-(4-benzyloxyphenyl)-1-cyclopentyl-N-methoxy-N-  
 methylbenzimidazole-5-carboxamide (Example 34),  
 30 2-(4-benzyloxyphenyl)-1-cyclopentyl-5-(1-hydroxy-1-  
 methylethyl)benzimidazole (Example 35),  
 5-acetyl-2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole  
 (Example 36),  
 2-(4-benzyloxyphenyl)-1-cyclopentyl-N-(2-dimethylaminoethyl)-  
 35 benzimidazole-5-carboxamide dihydrochloride (Example 37),  
 2-(4-benzyloxyphenyl)-1-cyclopentyl-5-nitrobenzimidazole  
 (Example 38),

- 5-amino-2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole hydrochloride (Example 39),
- 5-acetylamino-2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole (Example 40),
- 5 2-(4-benzyloxyphenyl)-1-cyclopentyl-5-methanesulfonyl-aminobenzimidazole (Example 41),
- 5-sulfamoyl-2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole (Example 42),
- 2-[4-(4-tert-butylbenzyloxy)phenyl]-1-cyclopentylbenzimidazole-10 5-carboxylic acid (Example 43),
- 2-[4-(4-carboxybenzyloxy)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 44),
- 2-[4-(4-chlorobenzyloxy)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 45),
- 15 2-[4-[(2-chloro-5-thienyl)methoxy]phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 46),
- 1-cyclopentyl-2-[4-(4-trifluoromethylbenzyloxy)phenyl]-benzimidazole-5-carboxylic acid (Example 47),
- 1-cyclopentyl-2-[4-(4-methoxybenzyloxy)phenyl]benzimidazole-5-20 carboxylic acid (Example 48),
- 1-cyclopentyl-2-[4-(4-pyridylmethoxy)phenyl]benzimidazole-5-carboxylic acid hydrochloride (Example 49),
- 1-cyclopentyl-2-[4-(4-methylbenzyloxy)phenyl]benzimidazole-5-carboxylic acid (Example 50),
- 25 1-cyclopentyl-2-[4-[(3,5-dimethyl-4-isoxazolyl)methoxy]phenyl]-benzimidazole-5-carboxylic acid (Example 51),
- 1-cyclopentyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylic acid (Example 52),
- [2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazol-5-yl]-30 carbonylaminoacetic acid (Example 53),
- 2-[4-(2-chlorobenzyloxy)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 54),
- 2-[4-(3-chlorobenzyloxy)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 55),
- 35 2-(4-benzyloxyphenyl)-3-cyclopentylbenzimidazole-5-carboxylic acid (Example 56),
- 2-[4-(benzenesulfonylamino)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 57),

- 1-cyclopentyl-2-[4-(3,5-dichlorophenylcarbonylamino)phenyl]-benzimidazole-5-carboxylic acid (Example 58),
- 2-[4-[(4-chlorophenyl)carbonylamino]phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 59),
- 5 2-[4-[(4-tert-butylphenyl)carbonylamino]phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 60),
- 2-[4-[(4-benzyloxyphenyl)carbonylamino]phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 61),
- trans-4-[2-(4-benzyloxyphenyl)-5-carboxybenzimidazol-1-yl]cyclohexan-1-ol (Example 62),
- 10 trans-1-[2-(4-benzyloxyphenyl)-5-carboxybenzimidazol-1-yl]-4-methoxycyclohexane (Example 63),
- 2-(4-benzyloxyphenyl)-5-carboxymethyl-1-cyclopentylbenzimidazole (Example 64),
- 15 2-[1-benzyloxycarbonyl-4-piperidyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 65),
- 2-[4-(cyclohexylphenyl)carbonylamino]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 66),
- 1-cyclopentyl-2-[4-(3,5-dichlorobenzyloxy)phenyl]benzimidazole-20 5-carboxylic acid (Example 67),
- 1-cyclopentyl-2-[4-(3,4-dichlorobenzyloxy)phenyl]benzimidazole-5-carboxylic acid (Example 68),
- 1-cyclopentyl-2-[4-(phenylcarbonylamino)phenyl]benzimidazole-5-carboxylic acid (Example 69),
- 25 1-cyclopentyl-2-[4-(diphenylmethoxy)phenyl]benzimidazole-5-carboxylic acid (Example 70),
- 1-cyclopentyl-2-(4-phenethyloxyphenyl)benzimidazole-5-carboxylic acid (Example 71),
- trans-1-[2-(4-benzyloxyphenyl)-5-carboxybenzimidazol-1-yl]-4-30 tert-butylcyclohexane (Example 72),
- 2-(4-benzyloxyphenyl)-5-carboxymethoxy-1-cyclopentylbenzimidazole (Example 73),
- 2-(4-benzylaminophenyl)-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 74),
- 35 2-[4-(N-benzenesulfonyl-N-methylamino)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 75),
- 2-[4-(N-benzyl-N-methylamino)phenyl]-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 76),

- 1-cyclohexyl-2-(4-phenethylphenyl)benzimidazole-5-carboxylic acid (Example 77),
- 2-(1-benzyl-4-piperidyl)-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 78),
- 5 2-(1-benzoyl-4-piperidyl)-1-cyclopentylbenzimidazole-5-carboxylic acid (Example 79),
- 1-cyclopentyl-2-[1-(p-toluenesulfonyl)-4-piperidyl]-benzimidazole-5-carboxylic acid (Example 80),
- 1-cyclohexyl-2-[4-(3,5-dichlorobenzyloxy)phenyl]benzimidazole-5-  
 10 carboxylic acid (Example 81),
- 1-cyclohexyl-2-[4-(diphenylmethoxy)phenyl]benzimidazole-5-carboxylic acid (Example 82),
- 1-cyclohexyl-2-[4-(3,5-di-tert-butylbenzyloxy)phenyl]-benzimidazole-5-carboxylic acid (Example 83),
- 15 2-(4-benzyloxyphenyl)-1-(4-methylcyclohexyl)benzimidazole-5-carboxylic acid (Example 84),
- 1-cyclohexyl-2-[4-[2-(2-naphthyl)ethoxy]phenyl]benzimidazole-5-carboxylic acid (Example 85),
- 1-cyclohexyl-2-[4-(1-naphthyl)methoxyphenyl]benzimidazole-5-  
 20 carboxylic acid (Example 86),
- 1-cyclohexyl-2-[4-(dibenzylamino)phenyl]benzimidazole-5-carboxylic acid (Example 87),
- 2-[4-(2-biphenyl)methoxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 88),
- 25 2-(4-benzyloxyphenyl)-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 89),
- 1-cyclohexyl-2-[4-(dibenzylmethoxy)phenyl]benzimidazole-5-carboxylic acid (Example 90),
- 2-(4-benzoylmethoxyphenyl)-1-cyclohexylbenzimidazole-5-  
 30 carboxylic acid (Example 91),
- 2-(4-benzyl-1-piperazinyl)-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 92),
- 1-cyclohexyl-2-[4-(3,3-diphenylpropyloxy)phenyl]benzimidazole-5-carboxylic acid (Example 93),
- 35 2-[4-(3-chloro-6-phenylbenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 94),
- 2-(4-benzyloxypiperidino)-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 95),

- 1-cyclohexyl-2-{4-[2-(phenoxy)ethoxy]phenyl}benzimidazole-5-carboxylic acid (Example 96),
- 1-cyclohexyl-2-[4-(3-phenylpropyloxy)phenyl]benzimidazole-5-carboxylic acid (Example 97),
- 5 1-cyclohexyl-2-[4-(5-phenylpentyloxy)phenyl]benzimidazole-5-carboxylic acid (Example 98),
- 2-(3-benzyloxy-5-isoxazolyl)-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 99),
- 2-(2-benzyloxy-5-pyridyl)-1-cyclohexylbenzimidazole-5-carboxylic
- 10 acid (Example 100),
- 1-cyclohexyl-2-{4-[2-(3,4,5-trimethoxyphenyl)ethoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 101),
- 2-(4-benzyloxyphenyl)-1-(4,4-dimethylcyclohexyl)benzimidazole-5-carboxylic acid (Example 102),
- 15 1-cyclohexyl-2-{4-[2-(1-naphthyl)ethoxy]phenyl}benzimidazole-5-carboxylic acid (Example 103),
- 2-[4-(2-benzyloxyphenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 104),
- 2-[4-(3-benzyloxyphenoxy)phenyl]-1-cyclohexylbenzimidazole-5-
- 20 carboxylic acid (Example 105),
- 1-cyclohexyl-2-[4-(2-hydroxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 106),
- 1-cyclohexyl-2-[4-(3-hydroxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 107),
- 25 1-cyclohexyl-2-[4-(2-methoxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 108),
- 1-cyclohexyl-2-[4-(3-methoxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 109),
- 1-cyclohexyl-2-[4-(2-propoxyphenoxy)phenyl]benzimidazole-5-
- 30 carboxylic acid (Example 110),
- 1-cyclohexyl-2-[4-(3-propoxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 111),
- 1-cyclohexyl-2-{4-[2-(3-methyl-2-butenyloxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 112),
- 35 1-cyclohexyl-2-{4-[3-(3-methyl-2-butenyloxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 113),
- 1-cyclohexyl-2-[4-(2-isopentyloxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 114),

- 1-cyclohexyl-2-[4-(3-isopentyloxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 115),
- 1-cyclohexyl-2-{4-[2-(10,11-dihydro-5H-dibenzo[b,f]azepin-5-yl)ethoxy]phenyl}benzimidazole-5-carboxylic acid (Example 116),
- 5 1-cyclohexyl-2-{4-[2-(4-trifluoromethylphenyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid (Example 117),
- 2-{4-[bis(4-chlorophenyl)methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 118),
- 1-cyclohexyl-2-{4-[2-(4-methoxyphenyl)ethoxy]phenyl}-
- 10 benzimidazole-5-carboxylic acid (Example 119),
- 1-cyclohexyl-2-{4-[2-(2-methoxyphenyl)ethoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 120),
- 1-cyclohexyl-2-{4-[2-(3-methoxyphenyl)ethoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 121),
- 15 2-(4-benzyloxyphenyl)-1-cycloheptylbenzimidazole-5-carboxylic acid (Example 122),
- 1-cyclohexyl-2-[4-(2-phenethyloxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 123),
- 1-cyclohexyl-2-[4-(3-phenethyloxyphenoxy)phenyl]benzimidazole-5-
- 20 carboxylic acid (Example 124),
- 1-cyclohexyl-2-[4-(2,2-diphenylethoxy)phenyl]benzimidazole-5-carboxylic acid (Example 125),
- 2-(4-benzyloxyphenyl)-1-(3-cyclohexenyl)benzimidazole-5-carboxylic acid (Example 126),
- 25 cis-1-[2-(4-benzyloxyphenyl)-5-carboxybenzimidazol-1-yl]-4-fluorocyclohexane (Example 127),
- 1-cyclohexyl-2-[4-(2-phenoxyphenoxy)phenyl]benzimidazole-5-carboxylic acid (Example 128),
- 1-cyclohexyl-2-[4-(3-phenoxyphenoxy)phenyl]benzimidazole-5-
- 30 carboxylic acid (Example 129),
- 2-{4-[(2R)-2-benzyloxycarbonylamino-2-phenylethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 130),
- 1-cyclohexyl-2-{2-fluoro-4-[2-(4-trifluoromethylphenyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid (Example 131),
- 35 2-[4-(4-benzyloxyphenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 132),
- 2-{4-[bis(4-methylphenyl)methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 133),



- 2-{4-[bis(4-fluorophenyl)methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 134),
- 1-cyclohexyl-6-methoxy-2-[4-(3-phenylpropoxy)phenyl]-benzimidazole-5-carboxylic acid (Example 135),
- 5 1-cyclohexyl-6-hydroxy-2-[4-(3-phenylpropoxy)phenyl]-benzimidazole-5-carboxylic acid (Example 136),
- 1-cyclohexyl-6-methyl-2-[4-(3-phenylpropoxy)phenyl]-benzimidazole-5-carboxylic acid (Example 137),
- 2-{4-[2-(2-benzyloxyphenyl)ethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 138),
- 10 2-{4-[2-(3-benzyloxyphenyl)ethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 139),
- 2-[4-(2-carboxymethyloxyphenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 140),
- 15 2-[4-(3-carboxymethyloxyphenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 141),
- 2-{4-[3-chloro-6-(4-methylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 142),
- 2-{4-[3-chloro-6-(4-methoxyphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 143),
- 20 1-cyclohexyl-2-{2-methyl-4-[2-(4-trifluoromethylphenyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid (Example 144),
- 2-{4-[2-(4-tert-butylphenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 145),
- 25 2-{4-(3-chloro-6-phenylbenzyloxy)-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 146),
- 2-{4-[3-chloro-6-(3,5-dichlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 147),
- 2-{4-[bis(4-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 148),
- 30 2-{4-(4-benzyloxyphenoxy)-2-chlorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 149),
- 2-{4-(4-benzyloxyphenoxy)-2-trifluoromethylphenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 150),
- 35 2-{4-[3-chloro-6-(2-trifluoromethylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 151),
- 2-{4-[(2R)-2-amino-2-phenylethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 152),

- 2-[4-(2-biphenyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 153),
- 2-[4-(3-biphenyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 154),
- 5 2-[4-[2-(1-tert-butoxycarbonyl-4-piperidyl)methoxy]phenoxy]-phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 155),
- 2-[4-[3-(1-tert-butoxycarbonyl-4-piperidyl)methoxy]phenoxy]-phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 156),
- 2-[4-[3-chloro-6-(3,4,5-trimethoxyphenyl)benzyloxy]phenyl]-1-
- 10 cyclohexylbenzimidazole-5-carboxylic acid (Example 157),
- 2-[4-[2-(2-biphenyl)ethoxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 158),
- 2-[4-(2-biphenylmethoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 159),
- 15 1-cyclohexyl-2-[4-[2-(4-piperidylmethoxy)phenoxy]phenyl]-benzimidazole-5-carboxylic acid hydrochloride (Example 160),
- 1-cyclohexyl-2-[4-[3-(4-piperidylmethoxy)phenoxy]phenyl]-benzimidazole-5-carboxylic acid hydrochloride (Example 161),
- 2-[4-[(2R)-2-acetylamino-2-phenylethoxy]phenyl]-1-
- 20 cyclohexylbenzimidazole-5-carboxylic acid (Example 162),
- 1-cyclohexyl-2-[4-[3-(4-methyl-3-pentenyl)phenoxy]phenyl]-benzimidazole-5-carboxylic acid (Example 163),
- 1-cyclohexyl-2-[4-[3-(3-methyl-3-butenyl)phenoxy]phenyl]-benzimidazole-5-carboxylic acid (Example 164),
- 25 2-[4-[(2S)-1-benzyl-2-pyrrolidinyl]methoxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 165),
- 2-[4-[3-chloro-6-(4-methylthiophenyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 166),
- 2-[4-[3-chloro-6-(4-methanesulfonylphenyl)benzyloxy]phenyl]-1-
- 30 cyclohexylbenzimidazole-5-carboxylic acid (Example 167),
- 2-[4-[3-chloro-6-(2-thienyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 168),
- 2-[4-[3-chloro-6-(3-chlorophenyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 169),
- 35 2-[4-[3-chloro-6-(3-pyridyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 170),
- 2-[4-[3-chloro-6-(4-fluorophenyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 171),

- 2-[4-(4-benzyloxyphenoxy)-3-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 172),
- 2-[4-(2-bromo-5-chlorobenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 173),
- 5 2-{4-[3-chloro-6-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 174),
- 2-{4-[2-{(1-acetyl-4-piperidyl)methoxy}phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 175),
- 2-{4-[3-{(1-acetyl-4-piperidyl)methoxy}phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 176),
- 10 1-cyclohexyl-2-{4-[3-(2-propynyloxy)phenoxy]phenyl}benzimidazole-5-carboxylic acid (Example 177),
- 1-cyclohexyl-2-{4-[3-(3-pyridylmethoxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 178),
- 15 2-(4-benzyloxy-2-methoxyphenyl)-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 179),
- 2-[4-(2-bromo-5-methoxybenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 180),
- 2-[4-(carboxydiphenylmethoxy)phenyl]-1-cyclohexylbenzimidazole-
- 20 5-carboxylic acid (Example 181),
- 2-{4-[2-(4-chlorophenyl)-5-nitrobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 182),
- 2-{4-[3-acetylamino-6-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 183),
- 25 2-{4-[2-(4-carboxyphenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 184),
- 2-{4-[(2S)-1-benzyloxycarbonyl-2-pyrrolidinyl]methoxy}phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 185),
- 2-{2-chloro-4-[2-(4-trifluoromethylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 186),
- 30 1-cyclohexyl-2-{4-[3-(2-pyridylmethoxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 187),
- 2-{4-[2-(4-chlorophenyl)-5-fluorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 188),
- 35 2-{4-[3-carboxy-6-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 189),
- 2-{4-[3-carbamoyl-6-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 190),

- 1-cyclohexyl-2-{4-[2-(dimethylcarbamoylmethoxy)phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 191),
- 1-cyclohexyl-2-{4-[2-(piperidinocarbonylmethoxy)phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 192),
- 5 2-{4-[(2S)-1-benzenesulfonyl-2-pyrrolidinyl]methoxy}phenyl-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 193),
- 2-{4-[(2S)-1-benzoyl-2-pyrrolidinyl]methoxy}phenyl-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 194),
- 2-{4-[2-(4-carbamoylphenyl)-5-chlorobenzyloxy]phenyl}-1-
- 10 cyclohexylbenzimidazole-5-carboxylic acid (Example 195),
- 1-cyclohexyl-2-{4-[3-(dimethylcarbamoylmethoxy)phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 196),
- 1-cyclohexyl-2-{4-[3-(piperidinocarbonylmethoxy)phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 197),
- 15 1-cyclohexyl-2-{4-[3-[(1-methanesulfonyl-4-piperidyl)methoxy]-phenoxy]phenyl}benzimidazole-5-carboxylic acid (Example 198),
- 1-cyclohexyl-2-{4-[(2-methyl-5-(4-chlorophenyl)-4-oxazolyl)-methoxy]phenyl}benzimidazole-5-carboxylic acid (Example 199),
- 2-{4-[3-(3-chlorobenzyloxy)phenoxy]phenyl}-1-
- 20 cyclohexylbenzimidazole-5-carboxylic acid (Example 200),
- 2-{4-[3-(4-chlorobenzyloxy)phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 201),
- 1-cyclohexyl-2-{4-[3-(4-fluorobenzyloxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 202),
- 25 1-cyclohexyl-2-{4-[(2S)-1-(4-nitrophenyl)-2-pyrrolidinyl]-methoxy}phenyl}benzimidazole-5-carboxylic acid (Example 203),
- 1-cyclohexyl-2-{4-[(2S)-1-phenyl-2-pyrrolidinyl]methoxy}-phenyl}benzimidazole-5-carboxylic acid hydrochloride (Example 204),
- 30 2-{4-[(2S)-1-(4-acetylamino-phenyl)-2-pyrrolidinyl]methoxy}-phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 205),
- 2-{4-[(5-(4-chlorophenyl)-2-methyl-4-thiazolyl)methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 206),
- 2-{4-[bis(3-fluorophenyl)methoxy]phenyl}-1-
- 35 cyclohexylbenzimidazole-5-carboxylic acid (Example 207),
- 1-cyclohexyl-2-{4-[2-(4-chlorophenyl)-3-nitrobenzyloxy]phenyl}-benzimidazole-5-carboxylic acid (Example 208),

- 1-cyclohexyl-2-{4-[3-(4-tetrahydropyranyloxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 209),
- 1-cyclohexyl-2-{4-[3-(4-trifluoromethylbenzyloxy)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 210),
- 5 1-cyclohexyl-2-{4-[3-{(1-methyl-4-piperidyl)methoxy}phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 211),
- 2-{4-[3-(4-tert-butylbenzyloxy)phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 212),
- 2-{4-[3-(2-chlorobenzyloxy)phenoxy]phenyl}-1-
- 10 cyclohexylbenzimidazole-5-carboxylic acid (Example 213),
- 1-cyclohexyl-2-{4-[3-(3-pyridyl)phenoxy]phenyl}benzimidazole-5-carboxylic acid (Example 214),
- 2-{4-[3-(4-chlorophenyl)phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 215),
- 15 1-cyclohexyl-2-{4-[3-(4-methoxyphenyl)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 216),
- 1-cyclohexyl-2-{4-[{4-(4-methanesulfonylphenyl)-2-methyl-5-thiazolyl}methoxy]phenyl}benzimidazole-5-carboxylic acid (Example 217),
- 20 2-{4-[{4-(4-chlorophenyl)-2-methyl-5-thiazolyl}methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 218),
- 2-{4-[1-(4-chlorobenzyl)-3-piperidyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 219),
- 1-cyclohexyl-2-{4-[3-{(2-methyl-4-thiazolyl)methoxy}phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 220),
- 25 1-cyclohexyl-2-{4-[3-{(2,4-dimethyl-5-thiazolyl)methoxy}phenoxy]-phenyl}benzimidazole-5-carboxylic acid (Example 221),
- 1-cyclohexyl-2-{4-[3-(3,5-dichlorophenyl)phenoxy]phenyl}-benzimidazole-5-carboxylic acid (Example 222),
- 30 2-{4-[1-(4-chlorobenzyl)-4-piperidyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 223),
- 2-{4-[3-(4-chlorobenzyloxy)piperidino]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 224),
- 2-{4-[4-carbamoyl-2-(4-chlorophenyl)benzyloxy]phenyl}-1-
- 35 cyclohexylbenzimidazole-5-carboxylic acid (Example 225),
- 2-{4-[4-(4-chlorobenzyloxy)piperidino]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 226),

- 2-{4-[3-{(2-chloro-4-pyridyl)methoxy}phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 227),
- 2-{4-[{(2S)-1-(4-dimethylcarbamoylphenyl)-2-pyrrolidinyl}-methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid  
5 (Example 228),
- 2-{4-[2-(4-chlorophenyl)-5-ethoxycarbonylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 229),
- 1-cyclohexyl-2-[4-(3-trifluoromethylphenoxy)phenyl]-benzimidazole-5-carboxylic acid (Example 230),
- 10 1-cyclohexyl-2-{4-[4-(4-dimethylcarbamoylphenyl)-2-methyl-5-thiazolyl}methoxy}phenyl}benzimidazole-5-carboxylic acid (Example 231),
- 2-{4-[2-(4-chlorophenyl)-5-dimethylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 232),
- 15 2-{4-[4-(4-chlorophenyl)-2-methyl-5-pyrimidinyl}methoxy}phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 233),
- 2-{4-[2-(4-chlorophenyl)-3-pyridyl}methoxy}phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride  
20 (Example 234),
- 2-{4-[3-(4-chlorophenyl)-2-pyridyl}methoxy}phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 235),
- 2-{4-[2-(3-chlorophenyl)-4-methylamino-1,3,5-triazin-6-yloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid  
25 trifluoroacetate (Example 236),
- 2-{4-[2-(4-chlorophenyl)-4-(5-tetrazolyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 237),
- 2-[4-(4-benzyloxy-6-pyrimidinyl)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 238),
- 30 1-cyclohexyl-2-{4-[4-(4-pyridylmethoxy)-6-pyrimidinyl]phenyl}-benzimidazole-5-carboxylic acid (Example 239),
- 2-{4-[4-(3-chlorophenyl)-6-pyrimidinyl]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 240),
- methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 241),  
35 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 242),

- ethyl 2-{4-[3-(4-chlorophenyl)pyridin-2-ylmethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 243),
- methyl 2-[4-(2-bromo-5-tert-butoxycarbonylbenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 244),
- 5 methyl 2-{4-[5-tert-butoxycarbonyl-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 245),
- methyl 2-{4-[5-carboxy-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate hydrochloride (Example 246),
- methyl 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 247),
- 10 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 248),
- 2-{4-[3-(tert-butylsulfamoyl)-6-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 249),
- 15 2-{4-[2-(4-chlorophenyl)-5-sulfamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid trifluoroacetate (Example 250),
- 2-(4-benzyloxycyclohexyl)-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 251),
- 20 2-[2-(2-biphenylyloxymethyl)-5-thienyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 252),
- 2-[2-(2-biphenylyloxymethyl)-5-furyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 253),
- 25 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-hydroxymethyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylic acid (Example 254),
- 1-cyclohexyl-2-{4-[4-(4-carboxyphenyl)-2-methyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylic acid hydrochloride (Example 255),
- 30 1-cyclohexyl-2-{2-fluoro-4-[4-fluoro-2-(3-fluorobenzoyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid (Example 256),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-sulfonic acid (Example 257),
- 35 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-3-cyclohexylbenzimidazole-4-carboxylic acid (Example 258),

- 1-cyclohexyl-2-{4-[3-dimethylcarbamoyl-5-(4-pyridylmethoxy)-phenoxy]phenyl}benzimidazole-5-carboxylic acid dihydrochloride (Example 259),
- 1-cyclohexyl-2-{4-[3-carboxy-5-(4-pyridylmethoxy)phenoxy]-phenyl}benzimidazole-5-carboxylic acid dihydrochloride (Example 260),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-4-carboxylic acid (Example 261),
- 2-{4-[3-carbamoyl-6-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 262),
- 2-{4-[(2-(4-carboxyphenyl)-3-pyridyl)methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 263),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-(4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 264),
- 2-{4-[2-(4-chlorophenyl)-5-dimethylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 265),
- 1-cyclohexyl-2-{4-[3-dimethylcarbamoyl-6-(4-trifluoromethylphenyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid hydrochloride (Example 266),
- 1-cyclohexyl-2-{4-[3-dimethylcarbamoyl-6-(4-methylthiophenyl)benzyloxy]phenyl}benzimidazole-5-carboxylic acid hydrochloride (Example 267),
- 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 268),
- 2-{4-[2-(4-chlorophenyl)-5-dimethylcarbamoylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 269),
- 2-{4-[3-carbamoyl-6-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 270),
- 2-{4-[3-dimethylcarbamoyl-6-(4-methanesulfonylphenyl)benzyloxy]-phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 271),



- 2-{4-[3-dimethylcarbamoyl-6-(3-pyridyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 272),
- 2-{4-[3-dimethylcarbamoyl-6-(4-dimethylcarbamoylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 273),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-(1-oxo-4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 274),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-(1,1-dioxo-4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 275),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]-2-fluorophenyl}-1-(4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 276),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]-2-fluorophenyl}-1-(1-oxo-4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 277),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]-2-fluorophenyl}-1-(1,1-dioxo-4-tetrahydrothiopyranyl)benzimidazole-5-carboxylic acid (Example 278),
- 2-{4-[2-(4-chlorophenyl)-5-dimethylsulfamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 279),
- 2-{4-[2-(4-chlorophenyl)-5-methanesulfonylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 280),
- methyl 2-{4-[5-carboxy-2-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate hydrochloride (Example 281),
- 2-{4-[2-(4-chlorophenyl)-5-dimethylaminobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 282),
- 2-{4-[2-(4-chlorophenyl)-5-methanesulfonylaminobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 283),

- 2-{4-[2-(4-chlorophenyl)-5-diethylcarbamoylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 284),
- 2-{4-[2-(4-chlorophenyl)-5-isopropylcarbamoylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 285),
- 2-{4-[2-(4-chlorophenyl)-5-piperidinocarbonylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 286),
- 2-{4-[2-(4-chlorophenyl)-5-(1-pyrrolidinyl) carbonylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 287),
- 2-{4-[2-(4-chlorophenyl)-5-(2-hydroxyethyl) carbamoylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 288),
- 2-{4-[2-(4-chlorophenyl)-5-(4-hydroxypiperidino)-carbonylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 289),
- 2-{4-[2-(4-chlorophenyl)-5-morpholinocarbonylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 290),
- 2-{4-[2-(4-chlorophenyl)-5-thiomorpholinocarbonylbenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 291),
- 2-{4-[3-(carboxymethylcarbamoyl)-6-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 292),
- 2-{4-[2-{4-(2-carboxyethyl)phenyl}-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 293),
- 2-{4-[3-chloro-6-(4-hydroxymethylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 294),
- 2-{4-[3-chloro-6-(4-methoxymethylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 295),
- 2-{4-[2-(3-carboxyphenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 296),

- 2-{4-[2-(4-chlorophenyl)-5-methylthiobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 297),
- 2-{4-[2-(4-chlorophenyl)-5-methylsulfinylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 298),
- 5 2-{4-[2-(4-chlorophenyl)-5-cyanobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 299),
- 2-{4-[bis(3-pyridyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 300),
- 2-{4-[bis(4-dimethylcarbamoylphenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 301),
- 10 sodium 2-{4-[2-thienyl-3-thienylmethoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 302),
- methyl 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example
- 15 303),
- sodium 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example 304),
- 2-{4-[5-carboxy-2-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 305),
- 20 2-{4-[2-(4-carboxyphenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 306),
- 2-{4-[2-(4-carbamoylphenyl)-5-(dimethylcarbamoyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 307),
- 25 2-{4-[5-amino-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 308),
- 2-{4-[5-(4-chlorophenyl)-2-methoxybenzylsulfinyl]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 309),
- 30 2-{4-[5-(4-chlorophenyl)-2-methoxybenzylsulfonyl]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 310),
- 2-{4-[2-(4-chlorophenyl)-5-methoxybenzylthio]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example
- 35 311),
- 2-{4-[bis(4-carboxyphenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 312),

2-[4-(phenyl-3-pyridylmethoxy)-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 313),  
 methyl 2-{4-[2-(4-chlorophenyl)-5-(methylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (Example  
 5 314),  
 2-{4-[5-chloro-2-(4-pyridyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 315),  
 2-{4-[2-(4-chlorophenyl)-5-(benzylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid  
 10 hydrochloride (Example 316),  
 2-{4-[2-(4-chlorophenyl)-5-(cyclohexylmethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid  
 hydrochloride (Example 317),  
 15 2-{4-[2-(4-chlorophenyl)-5-(4-pyridylmethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid  
 dihydrochloride (Example 318),  
 2-{4-[2-(4-chlorophenyl)-5-(N-benzyl-N-methylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic  
 20 acid hydrochloride (Example 319),  
 methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexyl-1H-indole-5-carboxylate (Example 501),  
 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexyl-1H-indole-5-carboxylic acid (Example 502),  
 25 2-(4-benzyloxyphenyl)-1-cyclopentyl-1H-indole-5-carboxylic acid (Example 503),  
 ethyl 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo[1,2-a]pyridine-7-carboxylate (Example 601),  
 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo[1,2-a]pyridine-7-carboxylic acid (Example 602), and  
 30 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-3-cyclohexyl-3H-imidazo[4,5-b]pyridine-6-carboxylic acid (Example 701).  
 (92) The fused ring compound of the formula [I] or a pharmaceutically acceptable salt thereof, which is selected from  
 35 the group consisting of  
 2-{4-[5-dimethylaminocarbonyl-2-(4-pyridyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 320),

- 2-{4-[2-(4-chlorophenyl)-5-(4-methylpiperazin-1-ylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 321),
- 2-{4-[2-(4-chlorophenyl)-5-{N-(3-pyridylmethyl) carbamoyl}-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 322),
- 2-{4-[2-(4-chlorophenyl)-5-{N-(2-pyridylmethyl) carbamoyl}-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 323),
- 10 2-{4-[2-(4-chlorophenyl)-5-(cyclohexylcarbamoyl)benzyloxy]-phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 324),
- 2-{4-[2-(4-chlorophenyl)-5-(2-pyridin-4-ylethylcarbamoyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 325),
- 15 2-{4-[(4-fluorophenyl){4-(dimethylaminocarbonyl)phenyl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 326),
- 2-{4-[(4-fluorophenyl)(4-carboxyphenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 327),
- 20 2-{4-[2-(4-chlorophenyl)-5-(4-oxopiperidinocarbonyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 328),
- 2-{4-[2-(4-chlorophenyl)-5-hydroxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 329),
- 25 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 330),
- 30 2-{4-[2-(4-chlorophenyl)-5-(N-isopropyl-N-methylcarbamoyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 331),
- 2-{4-[2-(4-chlorophenyl)-5-(phenylcarbamoyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 332),
- 35 2-{4-[2-(4-chlorophenyl)-5-(4-methoxypiperidinocarbonyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 333),

2-{4-[2-(4-chlorophenyl)-5-(3-hydroxypropyloxy)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 334), and  
2-{4-[2-(4-chlorophenyl)-5-(2-hydroxyethoxy)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example  
5 335).

(93) The fused ring compound of the formula [I] or a pharmaceutically acceptable salt thereof, which is selected from the group consisting of

methyl 2-[4-(2-bromo-5-nitrobenzyloxy)-2-fluorophenyl]-1-  
10 cyclohexylbenzimidazole-5-carboxylate (Example 336),

methyl 2-[4-{2-(4-chlorophenyl)-5-nitrobenzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 337),

methyl 2-[4-{5-amino-2-(4-chlorophenyl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example  
15 338),

methyl 2-[4-{2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (Example 339),

20 2-[4-{2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 340),

2-{4-[2-(4-chlorophenyl)-5-(4-methylpiperidin-1-ylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-  
25 carboxylic acid hydrochloride (Example 341),

2-{4-[5-acetyl-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 342),

2-{4-[2-(4-chlorophenyl)-5-{(4-hydroxypiperidin-1-ylcarbonyl)-methoxy}benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic  
30 acid (Example 343),

2-{4-[2-(4-chlorophenyl)-5-(2-methoxyethoxy)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 344),

35 2-{4-[2-(4-chlorophenyl)-5-{2-(2-methoxyethoxy)ethoxy}benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 345),

- 2-{4-[2-(4-chlorophenyl)-5-(isobutylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 346),
- 2-{4-[2-(4-chlorophenyl)-5-(2-methylthiazol-4-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 347),
- 5 2-{4-[2-(4-chlorophenyl)-5-(3,4-dihydroxypiperidin-1-ylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 348),
- 2-{4-[2-(4-chlorophenyl)-5-(3-methyl-1,2,4-oxadiazol-5-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid
- 10 hydrochloride (Example 349),
- 2-{4-[2-(4-chlorophenyl)-4-(isopropylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 350),
- 2-{4-[2-(4-chlorophenyl)-4-(piperidinocarbonyl)benzyloxy]phenyl}-
- 15 1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 351),
- 2-{4-[2-(4-chlorophenyl)-5-(1-hydroxy-2-methylpropan-2-yl)carbonyl]benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 352),
- 20 2-{4-[2-(4-chlorophenyl)-5-(4,4-dimethyl-2-oxazolin-2-yl)]benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 353),
- 2-{4-[2-(4-chlorophenyl)-4-(4-hydroxypiperidin-1-ylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-
- 25 carboxylic acid hydrochloride (Example 354),
- 2-{4-[2-(4-chlorophenyl)-4-(2-hydroxyethyl)carbonyl]benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 355),
- 2-{4-[2-(4-chlorophenyl)-4-(4-pyridylmethyl)carbonyl]benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid
- 30 (Example 356),
- 2-{4-[2-(4-chlorophenyl)-4-(dimethylcarbonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 357),
- 35 2-{4-[5-(2-aminothiazol-4-yl)-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 358),

- 2-{4-[2-(4-chlorophenyl)-5-(4-hydroxypiperidin-1-ylsulfonyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 359),
- 2-{4-[5-(dimethylcarbamoyl)-2-(4-fluorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 360),
- 2-{4-[5-(dimethylcarbamoyl)-2-(3-fluorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 361),
- 2-{4-[2-(5-chlorothiophen-2-yl)-5-(dimethylcarbamoyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 362),
- 2-{4-[2-bromo-5-(5-methyloxazol-2-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 363),
- 2-{4-[2-bromo-5-(5-methylthiazol-2-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 364),
- 2-{4-[2-(4-chlorophenyl)-5-(5-methyloxazol-2-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 365),
- 2-{4-[2-(4-chlorophenyl)-5-(5-methylthiazol-2-yl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 366),
- 2-{4-[2-(4-chlorophenyl)-5-tetrazol-5-ylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 367),
- 2-{4-[5-chloro-2-(4-cyanophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 368),
- 2-{4-[5-chloro-2-(4-tetrazol-5-ylphenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 369),
- 2-{4-[2-(4-chlorophenyl)-5-{2-(4-hydroxypiperidin-1-yl)ethoxy}benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 370),



- 2-{4-[2-(4-chlorophenyl)-5-(2-oxopiperidin-1-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 371),
- 2-{4-[3-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 372),
- 2-{4-[2-(4-chlorophenyl)-5-(N-hydroxyamidino)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 373),
- 10 2-{4-[2-(4-chlorophenyl)-5-(2,5-dihydro-5-oxo-4H-1,2,4-oxadiazol-3-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 374),
- 2-{4-[2-(4-chlorophenyl)-5-(2-oxo-3H-1,2,3,5-oxathiadiazol-4-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 375),
- 15 2-{4-[2-(4-chlorophenyl)-5-(2,5-dihydro-5-oxo-4H-1,2,4-thiadiazol-3-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 376),
- 20 2-{4-[2-(4-chlorophenyl)-5-(cyclopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 377),
- 2-{4-[2-(4-chlorophenyl)-5-(cyclobutylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 378),
- 25 2-{4-[2-(4-chlorophenyl)-5-(tert-butylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 379),
- 2-{4-[2-(4-chlorophenyl)-5-(isobutylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 380),
- 30 2-{4-[2-(4-chlorophenyl)-5-{(1-hydroxypropan-2-yl)carbamoyl}benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 381),
- 35 2-{4-[2-(4-chlorophenyl)-5-(methoxycarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 382),

- 2-{4-[2-(4-chlorophenyl)-5-{(2,3-dihydroxypropyl) carbamoyl}-benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 383),
- 2-{4-[2-(4-chlorophenyl)-5-(N-ethyl-N-methylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 384),
- 2-{4-[2-(4-chlorophenyl)-5-(N-methyl-N-propylcarbamoyl)-benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 385),
- 2-{4-[2-(4-chlorophenyl)-5-(N-isopropyl-N-methylcarbamoyl)-benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 386),
- 2-{4-[2-(4-chlorophenyl)-5-(2,6-dimethylpiperidin-1-ylcarbonyl)-benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 387),
- 2-{4-[5-(butylcarbamoyl)-2-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 388),
- 2-{4-[2-(4-chlorophenyl)-5-(propylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 389),
- 2-{4-[2-(4-chlorophenyl)-5-(ethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 390),
- 2-{4-[2-(4-chlorophenyl)-5-{(dimethylcarbamoyl) amino}benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 391),
- 2-{4-[2-(4-chlorophenyl)-5-{(morpholinocarbonyl) amino}benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 392),
- 2-{4-[2-(4-chlorophenyl)-5-ureidobenzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 393),
- 2-{4-[2-(4-chlorophenyl)-5-{(ethylcarbamoyl) amino}benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 394),

- 2-{4-[2-(4-chlorophenyl)-5-{(isopropylcarbamoyl)amino}benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 395),
- 2-{4-[2-(3,4-difluorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 396),
- 2-{4-[2-(2,4-difluorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 397),
- 2-{4-[2-(3,5-dichlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 398),
- 2-{4-[2-(3-chloro-4-fluorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 399),
- 2-{4-[2-(3,4-dichlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 400),
- 2-{4-[2-(4-chloro-2-fluorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 401),
- 2-{4-[2-(4-chloro-2-fluorophenyl)-5-(pyrrolidin-1-ylcarbonyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 402),
- 2-{4-[2-(4-chloro-3-fluorophenyl)-5-(pyrrolidin-1-ylcarbonyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 403),
- 2-{4-[2-(4-chloro-3-fluorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 404),
- 2-{4-[2-{4-(methylthio)phenyl}-5-(2-oxopyrrolidin-1-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 405),
- 2-{4-[2-{4-(methylthio)phenyl}-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 406),

- 2-{4-[4-chloro-2-(4-chlorophenyl)-5-(1,1-dioxoisothiazolidin-2-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 407),
- 2-{4-[4-chloro-2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 408),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylaminosulfonyl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 409),
- 10 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 410),
- 2-{4-[2-(4-chlorophenyl)-5-(4-hydroxypiperidin-1-ylcarbonyl)benzyloxy]-2-fluorophenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 411),
- 15 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 412),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]phenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 413),
- 20 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]phenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 414),
- 25 2-{4-[2-(4-chlorophenyl)-5-(4-hydroxypiperidin-1-ylcarbonyl)benzyloxy]phenyl}-1-cyclopentylbenzimidazole-5-carboxylic acid hydrochloride (Example 415),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]phenyl}-1-(tetrahydrothiopyran-4-yl)benzimidazole-5-carboxylic acid hydrochloride (Example 416),
- 30 2-{4-[2-(4-chlorophenyl)-5-(pyrrolidin-1-ylcarbonyl)benzyloxy]phenyl}-1-(tetrahydrothiopyran-4-yl)benzimidazole-5-carboxylic acid hydrochloride (Example 417),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-(tetrahydrothiopyran-4-yl)benzimidazole-5-carboxylic acid hydrochloride (Example 418),
- 35

- 2-{4-[2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy]-2-fluorophenyl}-1-(tetrahydrothiopyran-4-yl)benzimidazole-5-carboxylic acid hydrochloride (Example 419),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-piperidinobenzimidazole-5-carboxylic acid hydrochloride (Example 420),
- 2-{4-[2-(4-chlorophenyl)-5-(pyrrolidin-1-ylcarbonyl)benzyloxy]-2-fluorophenyl}-1-piperidinobenzimidazole-5-carboxylic acid (Example 421),
- 2-{4-[2-(4-chlorophenyl)-5-(2-imidazolin-2-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 422),
- 2-{4-[2-(4-chlorophenyl)-5-(2-oxooxazolidin-3-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 423),
- 2-{4-[2-(4-chlorophenyl)-5-(2-oxoimidazolidin-1-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 424),
- 2-{4-[2-(4-chlorophenyl)-5-(2-oxazolin-2-ylamino)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 425),
- 2-{4-[2-{2-[(dimethylcarbamoyl)methoxy]methyl}-4-(4-fluorophenyl)thiazol-5-yl}methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 426),
- 2-{4-[4-(4-fluorophenyl)-2-(4-hydroxypiperidin-1-ylmethyl)thiazol-5-yl}methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid dihydrochloride (Example 427),
- 2-{4-[4-(4-fluorophenyl)-2-[(carbamoylmethoxy)methyl]thiazol-5-yl}methoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 428),
- 2-{4-[4-(4-fluorophenyl)-2-(methylcarbamoyl)thiazol-5-yl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 429),
- 2-{4-[4-(4-fluorophenyl)-2-{(2-hydroxyethyl)carbamoyl}thiazol-5-yl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 430),

- 2-{4-[{2-(4-fluorophenyl)-5-(dimethylcarbamoyl)thiophen-3-yl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 431),
- 2-{4-[{2-(4-fluorophenyl)-5-(isopropylcarbamoyl)thiophen-3-yl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 432),
- 2-{4-[{2-(4-fluorophenyl)-5-(4-hydroxypiperidin-1-ylcarbonyl)thiophen-3-yl}methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 433),
- 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexyl-5-tetrazol-5-ylbenzimidazole (Example 434),
- 2-{4-[2-(4-carboxyphenyl)-5-chlorobenzyloxy]-2-fluorophenyl}-1-cyclohexyl-5-tetrazol-5-ylbenzimidazole hydrochloride (Example 435),
- 2-{4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]-2-fluorophenyl}-1-cyclohexyl-5-(2,5-dihydro-5-oxo-4H-1,2,4-oxadiazol-3-yl)benzimidazole hydrochloride (Example 436),
- 2-{4-[5-carboxy-2-(4-chlorophenyl)benzyloxy]-2-fluorophenyl}-5-cyano-1-cyclohexylbenzimidazole (Example 437),
- 2-{4-[2-(4-chlorophenyl)-5-(dimethylcarbamoyl)benzyloxy]-2-fluorophenyl}-5-cyano-1-cyclohexylbenzimidazole (Example 438),
- 2-{4-[{N-(4-dimethylcarbamoyl)-N-(4-fluorophenyl)amino}-methyl]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 439),
- 2-{5-[bis(3-fluorophenyl)methyl]-2-fluoro-4-hydroxyphenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 440),
- 2-{3-[bis(3-fluorophenyl)methyl]-2-fluoro-4-hydroxyphenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid (Example 441),
- 2-{4-[(3-dimethylcarbamoylphenyl)(4-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 442),
- 2-{4-[{3-(4-hydroxypiperidyl-1-ylcarbonyl)phenyl}(4-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 443),

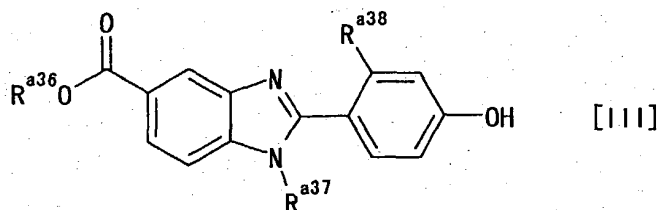
- 1-[2-[4-([4-(4-fluorophenyl)-2-methylthiazol-5-yl]methoxy)phenyl]-1-cyclohexylbenzimidazol-5-yl]carbonyl}-β-D-glucuronic acid (Example 444),
- {[2-[4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl]-1-cyclohexylbenzimidazol-5-yl]carbonyl}-β-D-glucuronic acid (Example 445),
- 2-[4-[2-(4-chlorophenyl)-5-(1,1-dioxoisothiazolidin-2-yl)benzyloxy]-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride (Example 446),
- 2-[4-[2-(4-chlorophenyl)-5-(isopropylcarbamoyl)benzyloxy]phenyl]-3-cyclohexyl-3H-dimidazo[4,5-b]pyridine-6-carboxylic acid hydrochloride (Example 702), and
- 2-[4-[2-(4-chlorophenyl)-5-(pyrrolidin-1-ylcarbonyl)benzyloxy]phenyl]-3-cyclohexyl-3H-imidazo[4,5-b]pyridine-6-carboxylic acid hydrochloride (Example 703).

- (94) A pharmaceutical composition comprising a fused ring compound of any of (42) to (93) above, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.
- (95) A hepatitis C virus polymerase inhibitor comprising a fused ring compound of any of (1) to (93) above, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.
- (96) An anti-hepatitis C virus agent comprising a fused ring compound of any of (1) to (93) above, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.
- (97) A therapeutic agent for hepatitis C comprising a fused ring compound of any of (42) to (93) above, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.
- (98) An anti-hepatitis C virus agent comprising (a) the anti-hepatitis C virus agent of (96) above and (b) at least one agent selected from the group consisting of a different antiviral agent, an antiinflammatory agent and an immunostimulant.
- (99) An anti-hepatitis C virus agent comprising (a) the anti-hepatitis C virus agent of (96) above and (b) interferon.

(100) A therapeutic agent for hepatitis C comprising (a) the hepatitis C virus polymerase inhibitor of (95) above and (b) at least one agent selected from the group consisting of a different antiviral agent, an antiinflammatory agent and an immunostimulant.

5 (101) A therapeutic agent for hepatitis C comprising (a) the hepatitis C virus polymerase inhibitor of (95) above and (b) interferon.

(102) A benzimidazole compound of the following formula [III]



10 wherein  $R^{a36}$  is hydrogen atom or carboxyl-protecting group,  $R^{a37}$  is cyclopentyl or cyclohexyl, and  $R^{a38}$  is hydrogen atom or fluorine atom, or a salt thereof.

(103) A thiazole compound selected from the group consisting of 4-(4-fluorophenyl)-5-hydroxymethyl-2-methylthiazole and 4-(4-  
15 fluorophenyl)-5-chloromethyl-2-methylthiazole, or a pharmaceutically acceptable salt thereof.

(104) A pharmaceutical composition comprising (a) the fused compound of the formula [I] of (1) above or a pharmaceutically acceptable salt thereof and (b) at least one agent selected from  
20 the group consisting of an antiviral agent other than the compound of (1) above, an antiinflammatory agent and an immunostimulant.

(105) A pharmaceutical composition comprising (a) the fused compound of the formula [I] of (1) above or a pharmaceutically  
25 acceptable salt thereof and (b) interferon.

(106) A method for treating hepatitis C, which comprises administering an effective amount of a fused ring compound of the formula [I] of (1) above, or a pharmaceutically acceptable salt thereof.

30 (107) The method of (106) above, further comprising administering an effective amount of at least one agent selected from the group consisting of an antiviral agent other than the compound of (1) above, an antiinflammatory agent and an immunostimulant.



(108) The method of (106) above, further comprising administering an effective amount of interferon.

(109) A method for inhibiting hepatitis C virus polymerase, which comprises administering an effective amount of a fused ring  
5 compound of the formula [I] of (1) above, or a pharmaceutically acceptable salt thereof.

(110) The method of (109) above, further comprising administering an effective amount of at least one agent selected from the group consisting of an antiviral agent other than the compound of (1)  
10 above, an antiinflammatory agent and an immunostimulant.

(111) The method of (109) above, further comprising administering an effective amount of interferon.

(112) Use of a fused ring compound of the above-mentioned formula [I] or a pharmaceutically acceptable salt thereof for the  
15 production of a pharmaceutical agent for treating hepatitis C.

(113) Use of a fused ring compound of the above-mentioned formula [I] or a pharmaceutically acceptable salt thereof for the production of a hepatitis C virus polymerase inhibitor.

(114) A pharmaceutical composition for the treatment of hepatitis  
20 C, which comprises a fused ring compound of the above-mentioned formula [I] or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

(115) A pharmaceutical composition for inhibiting hepatitis C virus polymerase, which comprises a fused ring compound of the  
25 above-mentioned formula [I] or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

(116) A commercial package comprising a pharmaceutical composition of (114) above and a written matter associated therewith, the written matter stating that the pharmaceutical  
30 composition can or should be used for treating hepatitis C.

(117) A commercial package comprising a pharmaceutical composition of (115) above and a written matter associated therewith, the written matter stating that the pharmaceutical composition can or should be used for inhibiting hepatitis C  
35 virus polymerase.

#### **Detailed Description of the Invention**

The definitions of respective substituents and moieties used in the present specification are as follows.

The halogen atom is a fluorine atom, chlorine atom, bromine atom or iodine atom, preferably fluorine atom, chlorine atom or bromine atom.

Particularly preferably, the halogen atom is fluorine atom  
5 at R<sup>5</sup>, R<sup>5'</sup>, R<sup>6</sup>, R<sup>6'</sup>, group A and group C, and fluorine atom or chlorine atom at X, Z, Z', group B and group D.

The C<sub>1-6</sub> alkyl is straight chain or branched chain alkyl having 1 to 6 carbon atoms, and is exemplified by methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl,  
10 isopentyl, tert-pentyl, hexyl and the like.

Preferably, it is straight chain or branched chain alkyl having 1 to 4 carbon atoms, and is particularly preferably methyl at R<sup>a7</sup>, R<sup>a8</sup>, R<sup>a9</sup>, R<sup>a15</sup>, R<sup>a16</sup>, R<sup>a17</sup>, R<sup>a29</sup>, R<sup>a33</sup>, R<sup>a35</sup>, R<sup>b6</sup> and R<sup>b7</sup> and methyl or tert-butyl at R<sup>b1</sup>, R<sup>b2</sup>, group B and group C.

15 The halogenated C<sub>1-6</sub> alkyl is the above-defined C<sub>1-6</sub> alkyl except that it is substituted by the above-defined halogen atom. Preferably, it is halogenated alkyl wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include fluoromethyl,  
20 difluoromethyl, trifluoromethyl, bromomethyl, chloromethyl, 1,2-dichloromethyl, 2,2-dichloromethyl, 2,2,2-trifluoroethyl and the like.

The halogenated C<sub>1-6</sub> alkyl is particularly preferably trifluoromethyl at group B.

25 The C<sub>1-6</sub> alkylene is straight chain alkylene having 1 to 6 carbon atoms, and is exemplified by methylene, ethylene, trimethylene, tetramethylene, pentamethylene or hexamethylene.

The C<sub>1-6</sub> alkylene is preferably methylene or ethylene at Y.

The C<sub>2-6</sub> alkenylene is straight chain alkenylene having 2  
30 to 6 carbon atoms, and is exemplified by vinylene, propenylene, 1-butenylene, 1,3-butadienylene and the like.

The C<sub>2-6</sub> alkenylene is preferably vinylene at Y.

The C<sub>1-6</sub> alkoxy is alkyloxy wherein the alkyl moiety thereof is the above-defined C<sub>1-6</sub> alkyl. Preferably, it is alkoxy  
35 wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutyloxy, tert-butyloxy, pentyloxy, hexyloxy and the like.

The C<sub>1-6</sub> alkoxy is particularly preferably methoxy at R<sup>a2</sup>, R<sup>a3</sup>, R<sup>a27</sup>, R<sup>a28</sup>, R<sup>a33</sup>, group A and group C.

The C<sub>1-6</sub> alkoxy C<sub>1-6</sub> alkoxy is that wherein C<sub>1-6</sub> alkoxy in the above definition is substituted by C<sub>1-6</sub> alkoxy defined above and  
5 is preferably that wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Specific examples include methoxymethyl, ethoxymethyl, methoxyethoxy, methoxypropoxy, isopropoxyethoxy and the like.

The group A is particularly preferably methoxyethoxy.

10 The C<sub>1-6</sub> alkanoyl is alkylcarbonyl wherein the alkyl moiety thereof is the above-defined C<sub>1-6</sub> alkyl. Preferably, it is alkanoyl wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include acetyl, propionyl, butyryl, isobutyryl, pivaloyl and the  
15 like.

The C<sub>1-6</sub> alkanoyl is particularly preferably acetyl at R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>a5</sup>, R<sup>a29</sup>, R<sup>b7</sup> and group B.

The C<sub>1-6</sub> alkoxycarbonyl is alkyloxycarbonyl wherein the alkoxy moiety thereof is the above-defined C<sub>1-6</sub> alkoxy. Preferably,  
20 it is alkoxycarbonyl wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxy carbonyl, butoxycarbonyl, isobutyloxycarbonyl, tert-butyloxycarbonyl, pentyloxycarbonyl,  
25 hexyloxycarbonyl and the like.

The C<sub>1-6</sub> alkoxycarbonyl is particularly preferably methoxycarbonyl or ethoxycarbonyl at R<sup>a10</sup> and group A.

The C<sub>1-6</sub> alkylamino is alkylamino or dialkylamino wherein the alkyl moiety thereof is the above-defined C<sub>1-6</sub> alkyl.  
30 Preferably, it is alkylamino or dialkylamino wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include methylamino, ethylamino, propylamino, isopropylamino, butylamino, isobutylamino, tert-butylamino, pentylamino, hexylamino,  
35 dimethylamino, diethylamino, methylethylamino, N-isopropyl-N-isobutylamino and the like.

The C<sub>1-6</sub> alkylamino is particularly preferably methylamino at R<sup>a7</sup>, and particularly preferably dimethylamino at R<sup>a21</sup> and group

A, and particularly preferably dimethylamino, ethylamino or isopropylamino at R<sup>a24</sup>.

The C<sub>1-6</sub> alkanoylamino is alkylcarbonylamino wherein the alkanoyl moiety thereof is the above-defined C<sub>1-6</sub> alkanoyl.

5 Preferably, it is alkylcarbonylamino wherein the alkyl moiety thereof is straight chain or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include acetylamino, propionylamino, butyrylamino, isobutyrylamino, pivaloylamino and the like.

10 The C<sub>1-6</sub> alkanoylamino is particularly preferably acetylamino at X and R<sup>a10</sup>.

The C<sub>1-6</sub> alkylsulfonyl is alkylsulfonyl wherein the alkyl moiety thereof is the above-defined C<sub>1-6</sub> alkyl. Preferably, it is alkylsulfonyl wherein the alkyl moiety thereof is straight chain  
15 or branched chain alkyl having 1 to 4 carbon atoms. Examples thereof include methylsulfonyl, ethylsulfonyl, propylsulfonyl, isopropylsulfonyl, butylsulfonyl, isobutylsulfonyl, tert-butylsulfonyl, pentylsulfonyl, hexylsulfonyl and the like.

The C<sub>1-6</sub> alkylsulfonyl is particularly preferably  
20 methylsulfonyl at X and R<sup>a5</sup>.

The C<sub>6-14</sub> aryl is aromatic hydrocarbon having 6 to 14 carbon atoms. Examples thereof include phenyl, naphthyl, anthryl, indenyl, azulenyl, fluorenyl, phenanthryl and the like.

The C<sub>6-14</sub> aryl is preferably phenyl or naphthyl,  
25 particularly preferably phenyl at the ring A, ring A', ring B and ring B'.

The C<sub>3-8</sub> cycloalkyl is saturated cycloalkyl having 3 to 8, preferably 5 to 7, carbon atoms. Examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and  
30 cyclooctyl.

The C<sub>3-8</sub> cycloalkyl is particularly preferably cyclohexyl at the ring A, ring A', ring B and ring B'.

The C<sub>3-8</sub> cycloalkenyl is cycloalkenyl having 3 to 8, preferably 5 to 7, carbon atoms and has at least 1, preferably 1  
35 or 2, double bond(s). Examples thereof include cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclopentadienyl, cyclohexenyl, 2,4-cyclohexadien-1-yl, 2,5-cyclohexadien-1-yl, cycloheptenyl and

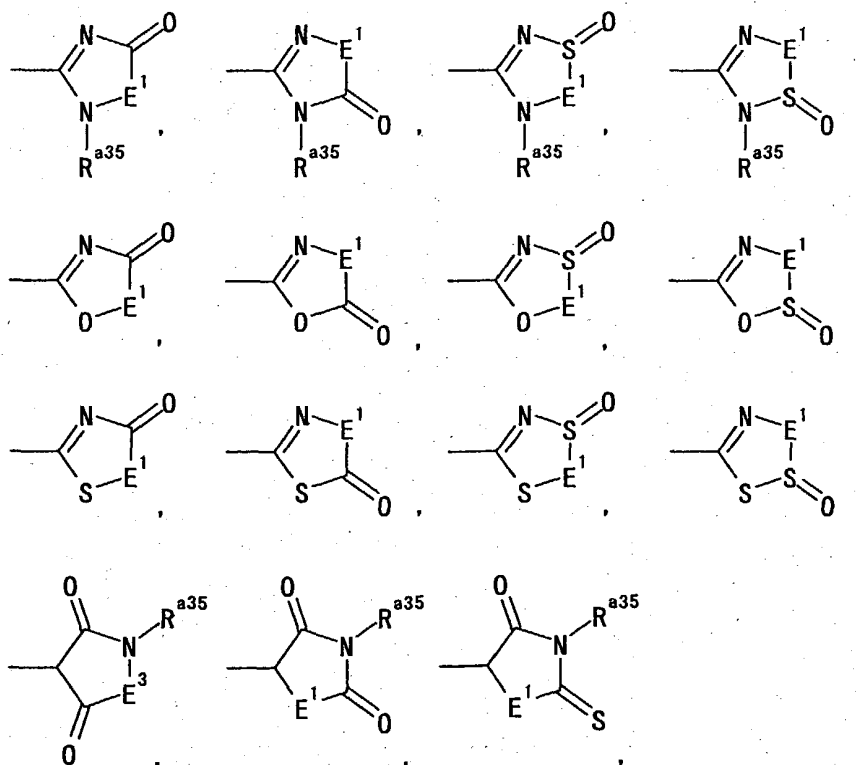
cyclooctenyl and the like, but do not include aryl (e.g., phenyl) or completely saturated cycloalkyl.

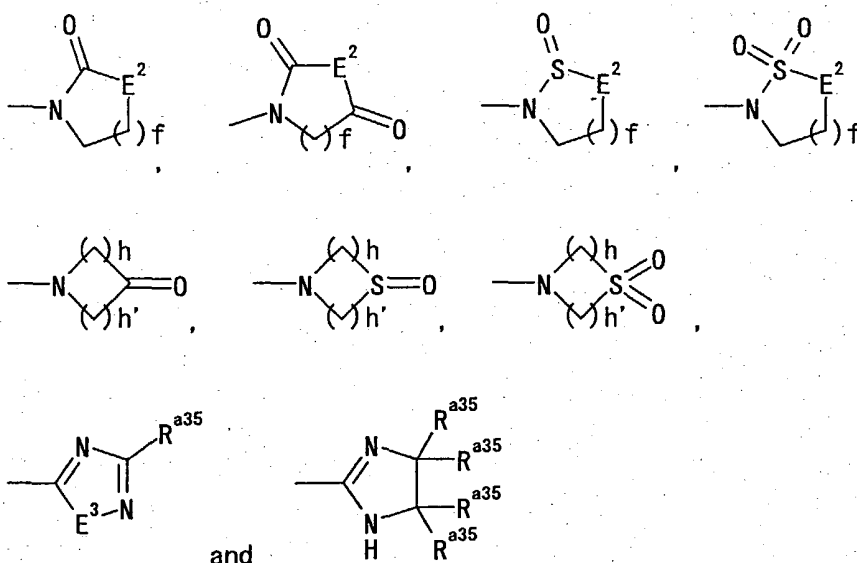
The C<sub>3-8</sub> cycloalkenyl is preferably cyclohexenyl at the ring A and ring A'.

- 5 The heterocyclic group has, as an atom constituting the ring, 1 to 4 heteroatom(s) selected from an oxygen atom, a nitrogen atom and a sulfur atom, besides a carbon atom, and includes saturated ring and unsaturated ring, monocyclic ring and fused ring having the number of ring atom constituting the ring  
10 of 3 to 14.

The heterocyclic group as a monocyclic ring includes, for example, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl,  
15 isothiazolyl, thiadiazolyl, pyrrolinyl, pyrrolidinyl, imidazolidinyl, piperidyl, piperazinyl, morpholinyl, thiomorpholinyl, tetrahydropyranyl and the like.

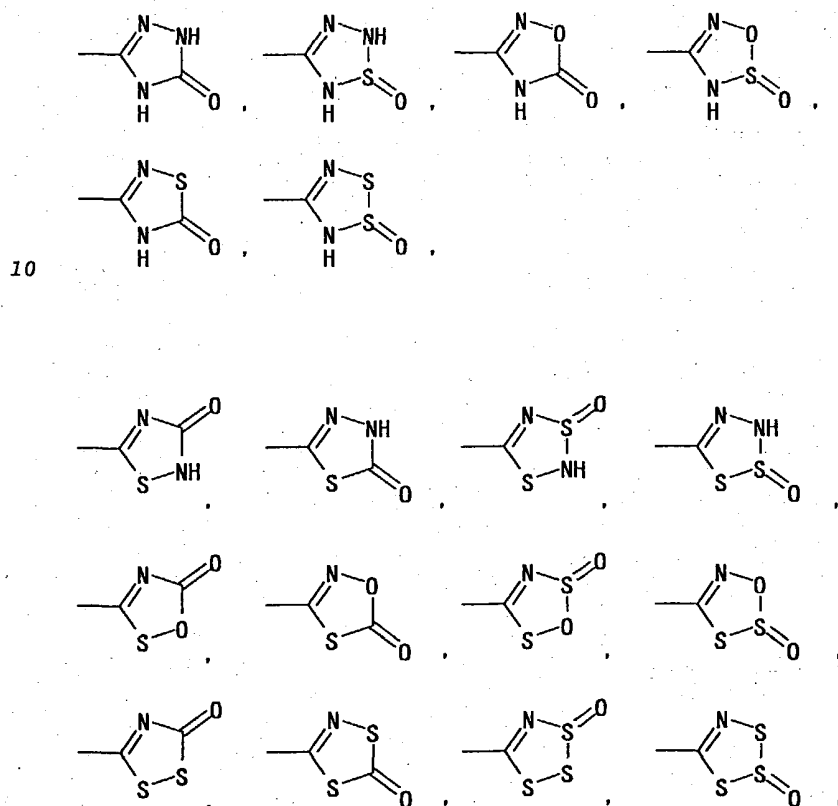
The heterocyclic group includes the groups of the following formulas.

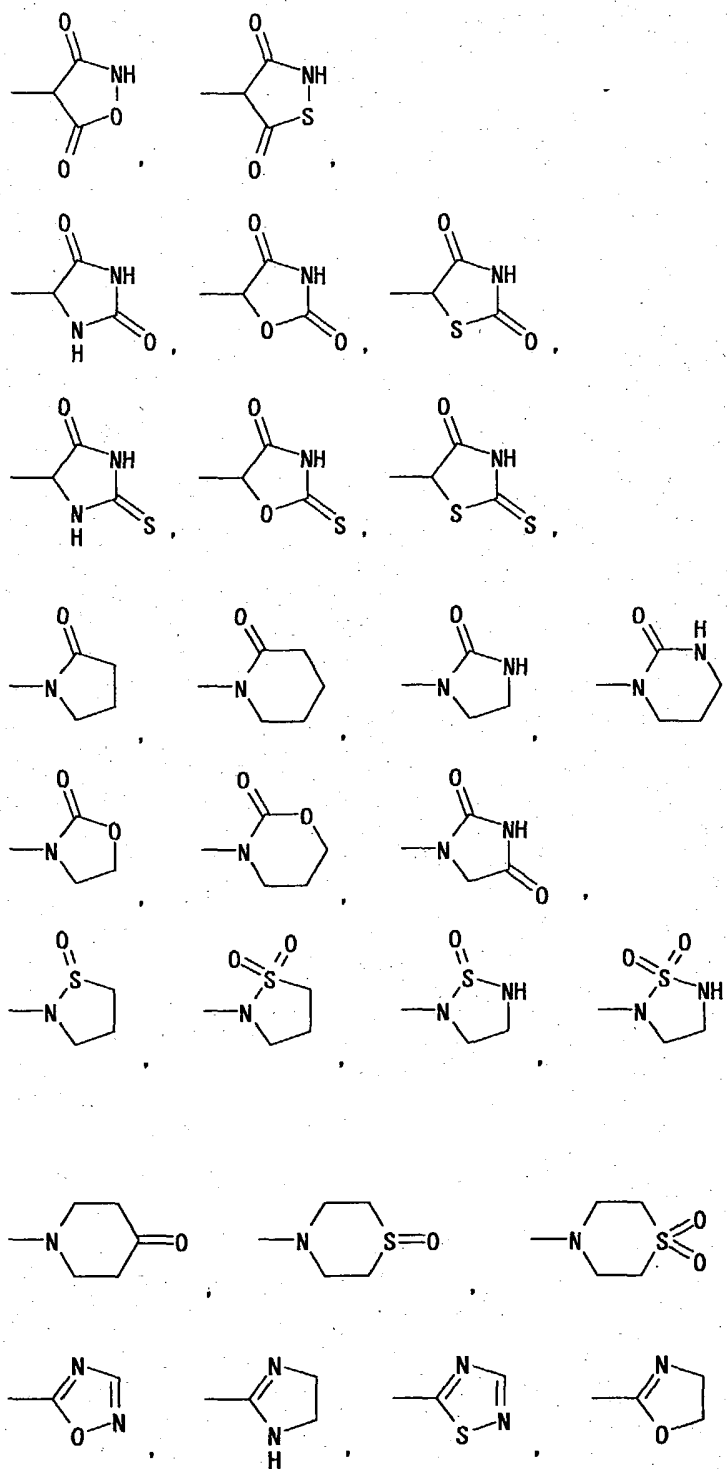




wherein  $E^1$  is an oxygen atom, a sulfur atom or  $N(-R^{a35})$ ,  $E^2$  is an  
 5 oxygen atom,  $CH_2$  or  $N(-R^{a35})$ ,  $E^3$  is an oxygen atom or a sulfur atom,  
 wherein  $R^{a35}$  is independently hydrogen atom or  $C_{1-6}$  alkyl,  $f$  is an  
 integer of 1 to 3, and  $h$  and  $h'$  are the same or different and  
 each is an integer of 1 to 3.

Specific examples of the heterocyclic group include



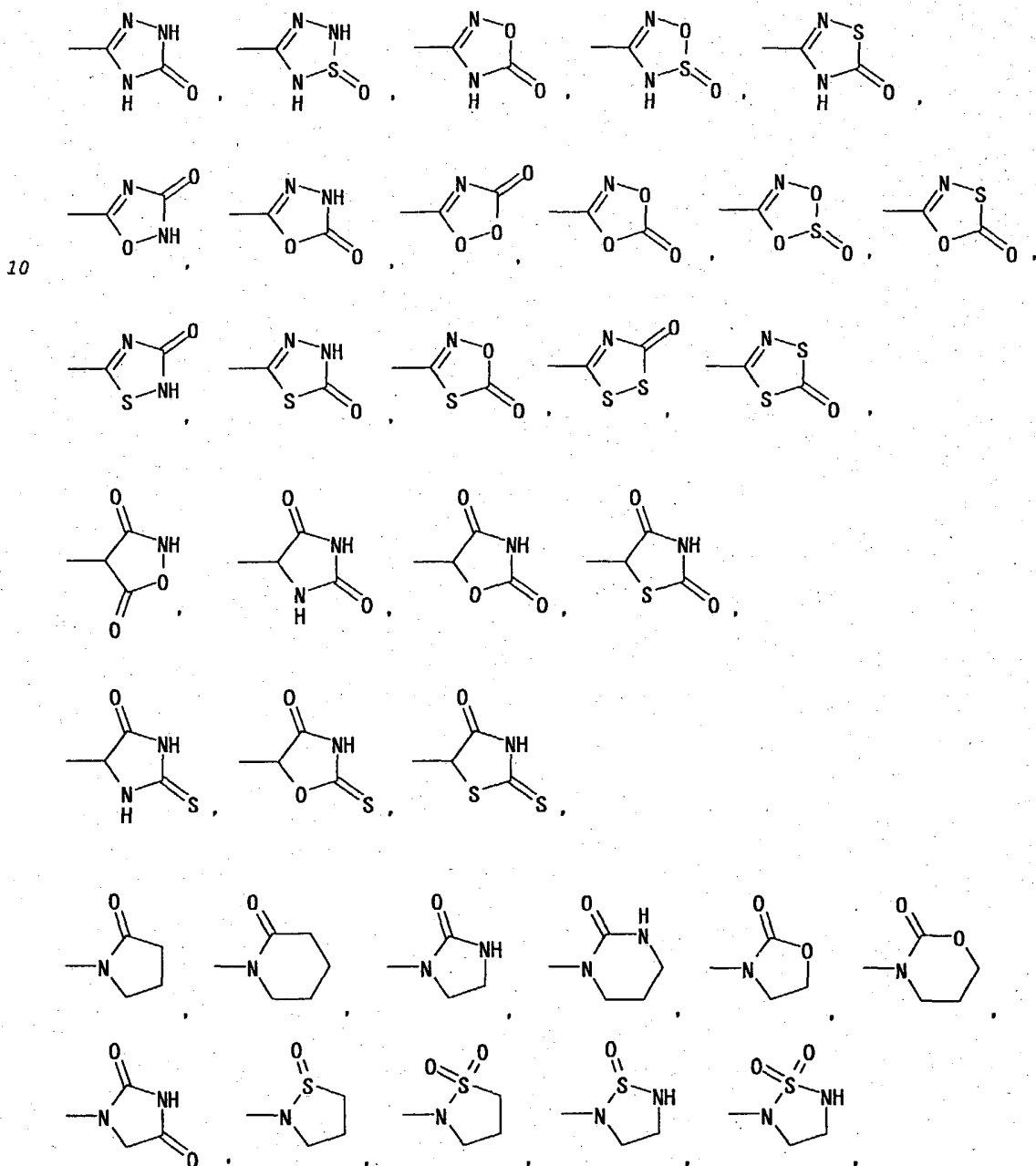


and the like.

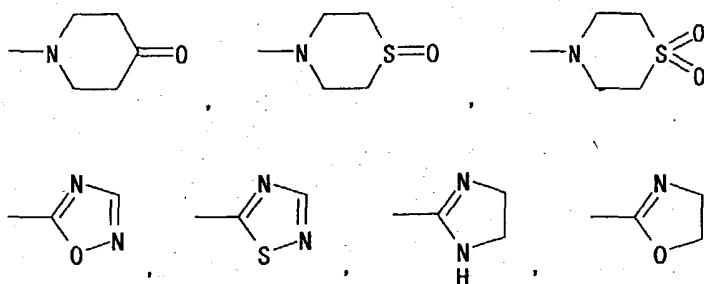
Examples of the heterocyclic group as a fused ring include quinolyl, isoquinolyl, quinazolinyl, quinoxalyl, phthalazinyl, cinnolinyl, naphthyridinyl, 5,6,7,8-tetrahydroquinolyl, indolyl,  
 10 benzimidazolyl, 2,3-dihydrobenzimidazolyl, 2,3-dihydro-2-

oxobenzimidazolyl, indolinyl, benzofuranyl, benzothienyl, benzoxazolyl, benzothiazolyl and the like.

Preferably, it is a heterocyclic group which is a 5-membered or a 6-membered monocyclic group. Examples thereof  
 5 include pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyrrolidinyl, piperidyl, piperazinyl







and the like.

At  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ ,  $Z$  and group D, tetrazolyl and 5-oxo- $\Delta^2$ -  
 5 1,2,4-oxadiazolin-3-yl are particularly preferable.

The heterocyclic group is preferably pyridyl, pyrazinyl, pyrimidinyl or pyridazinyl which is an aromatic group, and particularly preferably pyridyl at the ring A and ring A'.

The heterocyclic group is particularly preferably pyridyl,  
 10 pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl or thiadiazolyl, which is an aromatic group, at the ring B and ring B'. More preferably it is pyridyl or thiazolyl, most preferably  
 15 thiazolyl.

The  $C_{6-14}$  aryl  $C_{1-6}$  alkyl is arylalkyl wherein the alkyl moiety thereof is the above-defined  $C_{1-6}$  alkyl and the aryl moiety is the above-defined  $C_{6-14}$  aryl. Preferably, it is arylalkyl wherein the alkyl moiety thereof is straight chain alkyl having 1  
 20 to 4 carbon atoms and the aryl moiety is phenyl. Examples thereof include benzyl, phenethyl, 3-phenylpropyl, 2-phenylpropyl, 4-phenylbutyl and the like.

The  $C_{6-14}$  aryl  $C_{1-6}$  alkyl is particularly preferably benzyl at  $R^{a8}$  and  $R^{b6}$ .

25 The glucuronic acid residue is glucuronic acid less any hydroxyl group, preferably  $\beta$ -D-glucuronic acid substituted at 1-position.

The  $C_{6-14}$  aryl  $C_{1-6}$  alkyloxycarbonyl is arylalkyloxycarbonyl wherein the  $C_{6-14}$  aryl  $C_{1-6}$  alkyl moiety thereof is the above-  
 30 defined  $C_{6-14}$  aryl  $C_{1-6}$  alkyl. Preferably, it is arylalkyloxycarbonyl wherein the alkyl moiety thereof is straight chain alkyl having 1 to 4 carbon atoms and the aryl moiety is phenyl. Examples thereof include benzyloxycarbonyl,

phenethyloxycarbonyl, 3-phenylpropyloxycarbonyl, 2-phenylpropyloxycarbonyl, 4-phenylbutyloxycarbonyl and the like.

The C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyloxycarbonyl is particularly preferably benzyloxycarbonyl at R<sup>b7</sup>.

5        The optionally substituted C<sub>1-6</sub> alkyl is the above-defined C<sub>1-6</sub> alkyl, preferably that wherein straight chain or branched chain alkyl having 1 to 4 carbon atoms is optionally substituted with 1 to 3 substituent(s), and includes unsubstituted alkyl. The substituent(s) is(are) selected from the above-defined halogen  
10 atom, hydroxyl group, carboxyl, amino, the above-defined C<sub>1-6</sub> alkoxy, the above-defined C<sub>1-6</sub> alkoxy C<sub>1-6</sub> alkoxy, the above-defined C<sub>1-6</sub> alkoxy carbonyl and the above-defined C<sub>1-6</sub> alkylamino. Examples of optionally substituted C<sub>1-6</sub> alkyl include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl,  
15 pentyl, isopentyl, tert-pentyl, neopentyl, 1-ethylpropyl, hexyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, 3-hydroxypropyl, 4-hydroxybutyl, 1-hydroxy-1-methylethyl, 1-hydroxypropan-2-yl, 1,3-dihydroxypropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, carboxylmethyl, 2-carboxylethyl, methoxymethyl, methoxyethyl,  
20 methoxyethoxyethyl, ethoxycarbonylmethyl, 2-ethoxycarbonylethyl, 2-dimethylaminoethyl and the like.

Preferably, the optionally substituted C<sub>1-6</sub> alkyl is methyl, 1-hydroxy-1-methylethyl, carboxylmethyl or 2-dimethylaminoethyl at R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup>, methyl or trifluoromethyl at R<sup>5</sup>, R<sup>5'</sup>, R<sup>6</sup> and  
25 R<sup>6'</sup>, methyl at R<sup>7</sup>, R<sup>8</sup>, R<sup>a25</sup>, R<sup>a31</sup> and R<sup>b5</sup>, methyl, ethyl or isopropyl at R<sup>a24</sup>, methyl or isopropyl at R<sup>a18</sup>, methyl or ethyl at R<sup>a1</sup> and R<sup>a19</sup>, methyl, carboxylmethyl or 2-dimethylaminoethyl at R<sup>a2</sup> and R<sup>a3</sup>, methyl or carboxylmethyl at R<sup>a6</sup>, methyl, ethyl, isopropyl, butyl or trifluoromethyl at X, methyl, ethyl, isopropyl, butyl,  
30 isobutyl, tert-butyl, isopentyl, neopentyl, 1-ethylpropyl or carboxylmethyl at R<sup>a10</sup>, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, trifluoromethyl, 2-hydroxyethyl or carboxylmethyl at R<sup>a11</sup>, methyl or 4-hydroxybutyl at R<sup>a12</sup>, methyl, ethyl, isopropyl, butyl, 2-hydroxyethyl, 4-hydroxybutyl, ethoxycarbonylmethyl, 2-  
35 (ethoxycarbonyl)ethyl or 2-dimethylaminoethyl at R<sup>a13</sup>, methyl, propyl, butyl, isopentyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, 3-hydroxypropyl, methoxyethyl, methoxyethoxyethyl or carboxymethyl at R<sup>a20</sup>, methyl or ethyl at R<sup>a22</sup> and R<sup>a23</sup>, methyl

isopropyl or tert-butyl at  $R^{a26}$ , methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, isobutyl, 2-hydroxyethyl 1-hydroxypropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or carboxymethyl at  $R^{a27}$  and  $R^{a28}$ , and methyl, ethyl, propyl, isopropyl, tert-butyl, trifluoromethyl, 5 hydroxymethyl, 2-hydroxyethyl, 2-carboxylethyl, methoxymethyl or ethoxycarbonylmethyl at Z, Z' and group D.

It is particularly preferably, trifluoromethyl at  $R^5$ ,  $R^{5'}$ ,  $R^6$  and  $R^{6'}$ , methyl or tert-butyl at  $R^{a26}$ , methyl, tert-butyl, trifluoromethyl or hydroxymethyl at Z, Z' and group D, and methyl 10 at other substituents.

The optionally substituted  $C_{2-6}$  alkenyl is that wherein straight chain or branched chain alkenyl having 2 to 6 carbon atoms is optionally substituted by 1 to 3 substituent(s), and includes unsubstituted alkenyl. The substituent(s) is(are) 15 selected from the above-defined halogen atom, hydroxyl group, carboxyl, amino, the above-defined  $C_{1-6}$  alkoxy, the above-defined  $C_{1-6}$  alkoxy  $C_{1-6}$  alkoxy, the above-defined  $C_{1-6}$  alkoxycarbonyl and the above-defined  $C_{1-6}$  alkylamino. Examples of optionally substituted  $C_{2-6}$  alkenyl include vinyl, allyl, 1-propenyl, 20 isopropenyl, 1-butenyl, 2-butenyl, 1,3-butadienyl, 2-isopentenyl, 3-isohexenyl, 4-methyl-3-pentenyl, 2-carboxylethenyl and the like.

The optionally substituted  $C_{2-6}$  alkenyl is preferably 2-carboxylethenyl at X, and preferably 2-isopentenyl, 3-isohexenyl or 4-methyl-3-pentenyl at  $R^{a20}$ .

25 The optionally substituted  $C_{2-6}$  alkynyl is that wherein straight chain or branched chain alkynyl having 2 to 6 carbon atoms is optionally substituted by 1 to 3 substituent(s), and includes unsubstituted alkynyl. The substituent(s) is(are) selected from the above-defined halogen atom, hydroxyl group, 30 carboxyl, amino, the above-defined  $C_{1-6}$  alkoxy, the above-defined  $C_{1-6}$  alkoxycarbonyl and the above-defined  $C_{1-6}$  alkylamino. Examples thereof include ethynyl, 1-propynyl, 2-propynyl, 3-butynyl and the like.

The optionally substituted  $C_{2-6}$  alkynyl is preferably 2-propynyl at  $R^{a20}$ . 35

The  $C_{6-14}$  aryl optionally substituted by 1 to 5 substituent(s) selected from group B is that wherein the above-defined  $C_{6-14}$  aryl is optionally substituted by 1 to 5

substituent(s), and includes unsubstituted aryl. The substituent(s) is(are) selected from the above-defined halogen atom, cyano, nitro, the above-defined C<sub>1-6</sub> alkyl, the above-defined halogenated C<sub>1-6</sub> alkyl, the above-defined C<sub>1-6</sub> alkanoyl, - (CH<sub>2</sub>)<sub>r</sub>-COOR<sup>b1</sup>, - (CH<sub>2</sub>)<sub>r</sub>-CONR<sup>b1</sup>R<sup>b2</sup>, - (CH<sub>2</sub>)<sub>r</sub>-NR<sup>b1</sup>R<sup>b2</sup>, - (CH<sub>2</sub>)<sub>r</sub>-NR<sup>b1</sup>-COR<sup>b2</sup>,  
 5 - (CH<sub>2</sub>)<sub>r</sub>-NHSO<sub>2</sub>R<sup>b1</sup>, - (CH<sub>2</sub>)<sub>r</sub>-OR<sup>b1</sup>, - (CH<sub>2</sub>)<sub>r</sub>-SR<sup>b1</sup>, - (CH<sub>2</sub>)<sub>r</sub>-SO<sub>2</sub>R<sup>b1</sup> and - (CH<sub>2</sub>)<sub>r</sub>-SO<sub>2</sub>NR<sup>b1</sup>R<sup>b2</sup> (wherein R<sup>b1</sup> and R<sup>b2</sup> are each independently hydrogen atom or the above-defined C<sub>1-6</sub> alkyl and r is 0 or an integer of 1 to 6).

10 Examples thereof include phenyl, naphthyl, anthryl, indenyl, azulenyl, fluorenyl, phenanthryl, 3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 2,4-dichlorophenyl, 3,5-dichlorophenyl, pentafluorophenyl, 4-methylphenyl, 4-tert-butylphenyl, 2-trifluoromethylphenyl, 4-trifluoromethylphenyl, 4-  
 15 nitrophenyl, 4-cyanophenyl, 4-acetylphenyl, 4-carboxylphenyl, 4-carbamoylphenyl, 4-aminophenyl, 4-dimethylaminophenyl, 4-acetylaminophenyl, 4-(methylsulfonylamino)phenyl, 4-methoxyphenyl, 3,4,5-trimethoxyphenyl, 4-methylthiophenyl, 4-methylsulfonylphenyl, 4-aminosulfonylphenyl, 3-nitro-4-  
 20 methoxyphenyl and 4-nitro-3-methoxyphenyl.

The aryl moiety is preferably phenyl, the group B here is preferably the above-defined halogen atom, nitro, the above-defined C<sub>1-6</sub> alkyl, the above-defined halogenated C<sub>1-6</sub> alkyl or - (CH<sub>2</sub>)<sub>r</sub>-OR<sup>b1</sup>. Examples of group B include fluorine atom, chlorine  
 25 atom, nitro, methyl, tert-butyl, trifluoromethyl and methoxy. Particularly preferably, it is fluorine atom or chlorine atom.

With regard to "C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group B", it is preferably phenyl, 4-tert-butylphenyl, 4-fluorophenyl, 3-chlorophenyl, 4-  
 30 chlorophenyl, 4-methoxyphenyl or 4-trifluoromethylphenyl at R<sup>a12</sup>, R<sup>a27</sup> and R<sup>a28</sup>, phenyl at R<sup>a14</sup>, R<sup>a22</sup>, R<sup>a23</sup>, R<sup>a26</sup> and R<sup>b5</sup>, phenyl or 3-fluorophenyl at R<sup>a18</sup>, phenyl or 2,4-dichlorophenyl at R<sup>a20</sup>, phenyl, 4-chlorophenyl, 4-trifluoromethylphenyl, 3,5-dichlorophenyl, 3-nitro-4-methoxyphenyl or 4-nitro-3-methoxyphenyl at R<sup>a24</sup>, and  
 35 phenyl or 4-methylphenyl at R<sup>a25</sup>.

It is particularly preferably phenyl at other substituents.

The C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D is that wherein the above-

defined C<sub>6-14</sub> aryl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted aryl. The substituent(s) is(are) selected from the above-mentioned group D (substituents shown under (a) to (q)).

5        Examples of group D here include fluorine atom, chlorine atom, bromine atom, nitro, cyano, methyl, ethyl, propyl, isopropyl, tert-butyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, methoxymethyl, 2-carboxylethyl, methoxycarbonylmethyl, ethoxycarbonylmethyl, acetyl, carboxyl,  
10 methoxycarbonyl, ethoxycarbonyl, carbamoyl, methylaminocarbonyl, isopropylaminocarbonyl, dimethylaminocarbonyl, diethylaminocarbonyl, (2-hydroxyethyl)aminocarbonyl, (carboxylmethyl)aminocarbonyl, hydroxyl group, methoxy, ethoxy, propyloxy, isopropyloxy, isopentyloxy, 2-isopentyloxy, 3-  
15 isohexenyloxy, 4-methyl-3-pentyloxy, 2-propynyloxy, hydroxymethyloxy, carboxylmethyloxy, (dimethylaminocarbonyl)methyloxy, amino, methylamino, dimethylamino, diethylamino, acetylamino, methylsulfonylamino, methylthio, methylsulfonyl, methylsulfinyl, aminosulfonyl,  
20 methylaminosulfonyl, dimethylaminosulfonyl and tetrazolyl.

      Examples of C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D include phenyl, naphthyl, anthryl, indenyl, azulenyl, fluorenyl, phenanthryl, 3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl,  
25 2,4-dichlorophenyl, 3,5-dichlorophenyl, 4-bromophenyl, 4-nitrophenyl, pentafluorophenyl, 4-methylphenyl, 4-tert-butylphenyl, 2-trifluoromethylphenyl, 4-trifluoromethylphenyl, 4-(hydroxymethyl)phenyl, 4-(methoxymethyl)phenyl, 4-(2-carboxylethyl)phenyl, 3-carboxylphenyl, 4-carboxylphenyl, 4-  
30 methoxyphenyl, 3,4,5-trimethoxyphenyl, 4-carbamoylphenyl, 4-methylthiophenyl, 4-(dimethylaminocarbonyl)phenyl, 4-methylsulfonylphenyl, 4-acetylamino, 4-cyanophenyl, 4-acetylphenyl, 4-aminophenyl, 4-dimethylaminophenyl, 4-(methylsulfonylamino)phenyl, 4-methylsulfinylphenyl, 4-  
35 aminosulfonylphenyl and 3-nitro-4-methoxyphenyl, 4-nitro-3-methoxyphenyl and 4-tetrazol-5-ylphenyl.

      At Z and Z', the aryl moiety is preferably phenyl, and group D here is preferably the above-defined halogen atom, nitro,

the above-defined optionally substituted  $C_{1-6}$  alkyl,  $-(CH_2)_t-COOR^{a19}$ ,  
-  $(CH_2)_t-CONR^{a27}R^{a28}$ , -  $(CH_2)_t-OR^{a20}$ , -  $(CH_2)_t-NR^{a29}CO-R^{a24}$ ,  
-  $(CH_2)_t-S(O)_q-R^{a25}$  or -  $(CH_2)_t-SO_2-NHR^{a26}$ .

Examples of  $C_{6-14}$  aryl optionally substituted by 1 to 5  
5 substituent(s) selected from group D preferably include phenyl,  
3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl,  
3,5-dichlorophenyl, 4-bromophenyl, 4-nitrophenyl, 4-methylphenyl,  
4-tert-butylphenyl, 2-trifluoromethylphenyl, 4-  
trifluoromethylphenyl, 4-(hydroxymethyl)phenyl, 4-  
10 (methoxymethyl)phenyl, 4-(2-carboxylethyl)phenyl, 3-  
carboxylphenyl, 4-carboxylphenyl, 4-methoxyphenyl, 3,4,5-  
trimethoxyphenyl, 4-carbamoylphenyl, 4-methylthiophenyl, 4-  
(dimethylaminocarbonyl)phenyl, 4-methylsulfonylphenyl, 4-  
acetylaminophenyl, 4-methylsulfinylphenyl, 4-aminosulfonylphenyl,  
15 4-cyanophenyl and 4-tetrazolylphenyl.

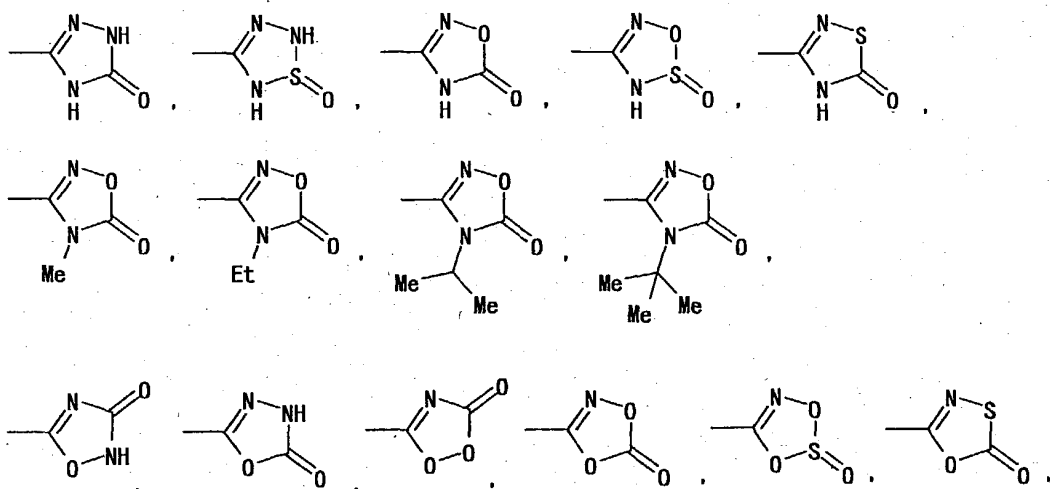
Particularly preferably, it is the above-defined halogen  
atom, the above-defined optionally substituted  $C_{1-6}$  alkyl,  
-  $(CH_2)_t-COOR^{a19}$ , -  $(CH_2)_t-CONR^{a27}R^{a28}$ , -  $(CH_2)_t-OR^{a20}$  or -  $(CH_2)_t-S(O)_q-R^{a25}$ ,  
which is specifically fluorine atom, chlorine atom, bromine atom,  
20 nitro, methyl, tert-butyl, carboxyl, trifluoromethyl,  
hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl,  
methylthio, dimethylaminocarbonyl, methylsulfonyl or acetylamino.  
More preferably, it is fluorine atom, chlorine atom, methyl,  
tert-butyl, carboxyl, methoxy, carbamoyl, methylthio,  
25 dimethylaminocarbonyl, methylsulfonyl or acetylamino, most  
preferably fluorine atom or chlorine atom.

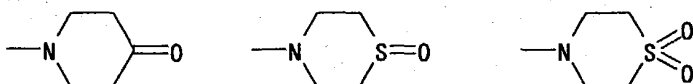
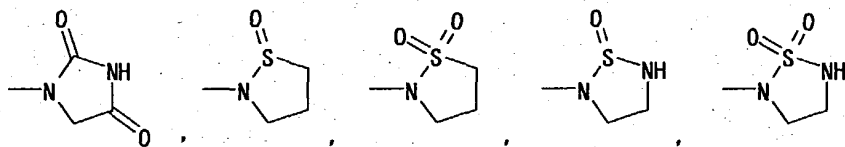
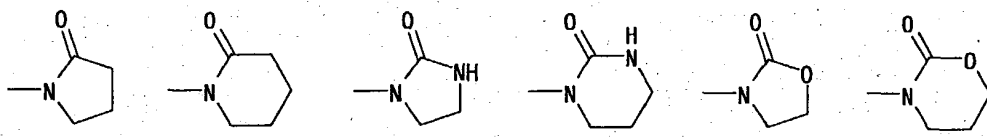
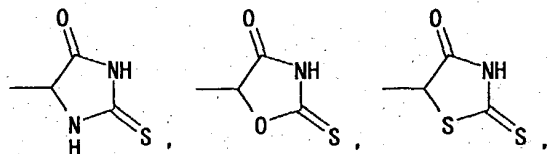
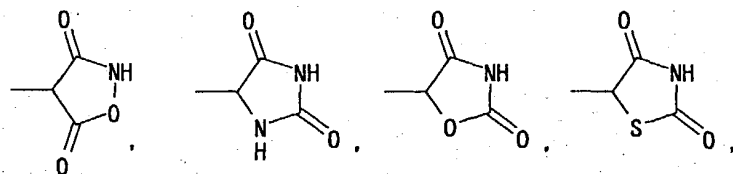
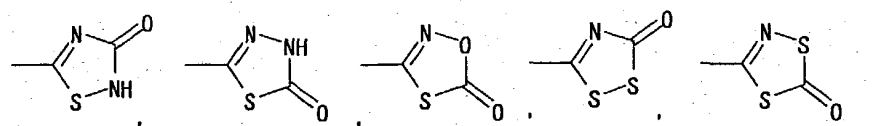
The heterocyclic group optionally substituted by 1 to 5  
substituent(s) selected from group B is that wherein the above-  
defined heterocyclic group is optionally substituted by 1 to 5  
30 substituent(s), and includes unsubstituted heterocyclic group.  
The substituent(s) is(are) selected from the above-defined  
halogen atom, cyano, nitro, the above-defined  $C_{1-6}$  alkyl, the  
above-defined halogenated  $C_{1-6}$  alkyl, the above-defined  $C_{1-6}$   
alkanoyl,

35 -  $(CH_2)_r-COOR^{b1}$ , -  $(CH_2)_r-CONR^{b1}R^{b2}$ , -  $(CH_2)_r-NR^{b1}R^{b2}$ , -  $(CH_2)_r-NR^{b1}-COR^{b2}$ ,  
-  $(CH_2)_r-NHSO_2R^{b1}$ , -  $(CH_2)_r-OR^{b1}$ , -  $(CH_2)_r-SR^{b1}$ , -  $(CH_2)_r-SO_2R^{b1}$  and

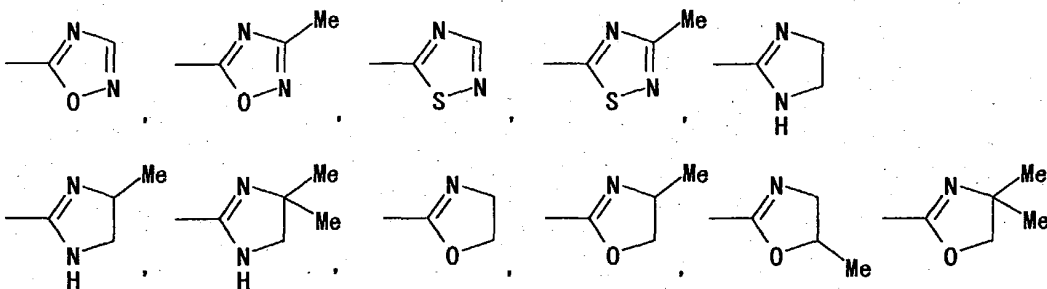
$-(CH_2)_r-SO_2NR^{b1}R^{b2}$  wherein  $R^{b1}$  and  $R^{b2}$  are each independently hydrogen atom or the above-defined  $C_{1-6}$  alkyl and  $r$  is 0 or an integer of 1 to 6.

Examples thereof include 2-pyridyl, 3-pyridyl, 4-pyridyl,  
 5 3-fluoropyridin-4-yl, 3-chloropyridin-4-yl, 4-chloropyridin-3-yl,  
 pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl,  
 pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, 2-thienyl, 3-  
 thienyl, furyl, oxazolyl, 2-methyloxazol-4-yl, isoxazolyl,  
 thiazolyl, 2-methylthiazol-4-yl, 2,5-dimethylthiazol-4-yl, 2,4-  
 10 dimethylthiazol-5-yl, isothiazolyl, thiadiazolyl, pyrrolinyl,  
 pyrrolidinyl, 3-hydroxypyrrolidinyl, imidazolidinyl, azetidiny,   
 piperidyl, 3-hydroxypiperidino, 4-hydroxypiperidino, 3,4-  
 dihydroxypiperidino, 4-methoxypiperidino, 4-carboxypiperidino, 4-  
 (hydroxymethyl)piperidino, 2-oxopiperidino, 4-oxopiperidino,  
 15 2,2,6,6-tetramethylpiperidino, 2,2,6,6-tetramethyl-4-  
 hydroxypiperidino, N-methylpiperidin-4-yl, N-(tert-  
 butoxycarbonyl)piperidin-4-yl, N-acetylpiperidin-4-yl, N-  
 methylsulfonylpiperidin-4-yl, piperazinyl, 4-methylpiperazinyl,  
 4-methylsulfonylpiperazinyl, morpholinyl, thiomorpholinyl, 1-  
 20 oxothiomorpholin-4-yl, 1,1-dioxothiomorpholin-4-yl,  
 tetrahydropyranyl, quinolyl, isoquinolyl, quinazolinyl,  
 quinoxalyl, phthalazinyl, cinnolinyl, naphthyridinyl, 5,6,7,8-  
 tetrahydroquinolyl, indolyl, benzimidazolyl, indolinyl,  
 benzofuranyl, benzothienyl, benzoxazolyl, benzothiazolyl,





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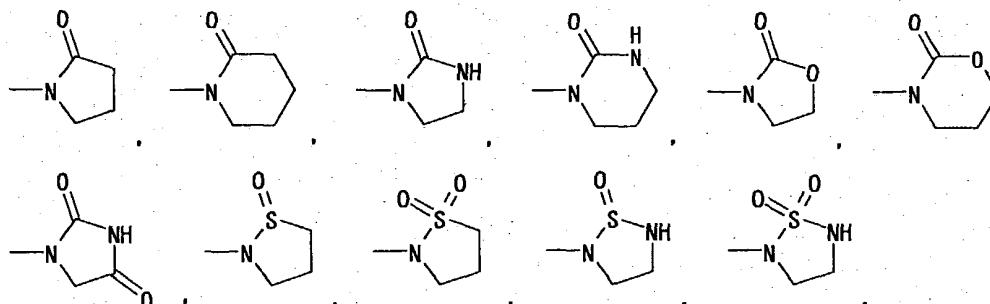
and the like.

The heterocyclic moiety is preferably a heterocyclic group which is a 5-membered or a 6-membered monocyclic group. Examples thereof include pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl,

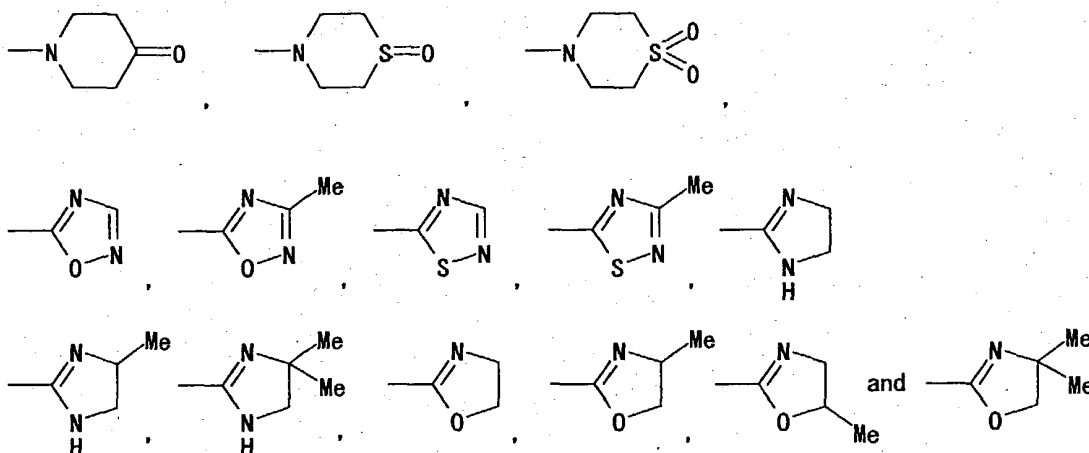


tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyrrolidinyl, piperidyl, piperazinyl, morpholinyl, thiomorpholinyl and tetrahydropyranyl, and the group B here is preferably the above-defined halogen atom, the above-defined  
 5 C<sub>1-6</sub> alkyl, the above-defined halogenated C<sub>1-6</sub> alkyl, the above-defined C<sub>1-6</sub> alkanoyl,  $-(CH_2)_r-COOR^{b1}$ ,  $-(CH_2)_r-CONR^{b1}R^{b2}$  or  $-(CH_2)_r-OR^{b1}$ .

Examples of heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from group B preferably include piperidino, 4-methylpiperidino, 2,6-dimethylpiperidino, 4-  
 10 hydroxypiperidino, 1-piperazinyl, 1-(methylsulfonyl)piperidin-4-yl, 1-pyrrolidinyl, morpholino, 4-thiomorpholinyl, tetrahydropyranyl, pyridyl, thiazolyl,



15



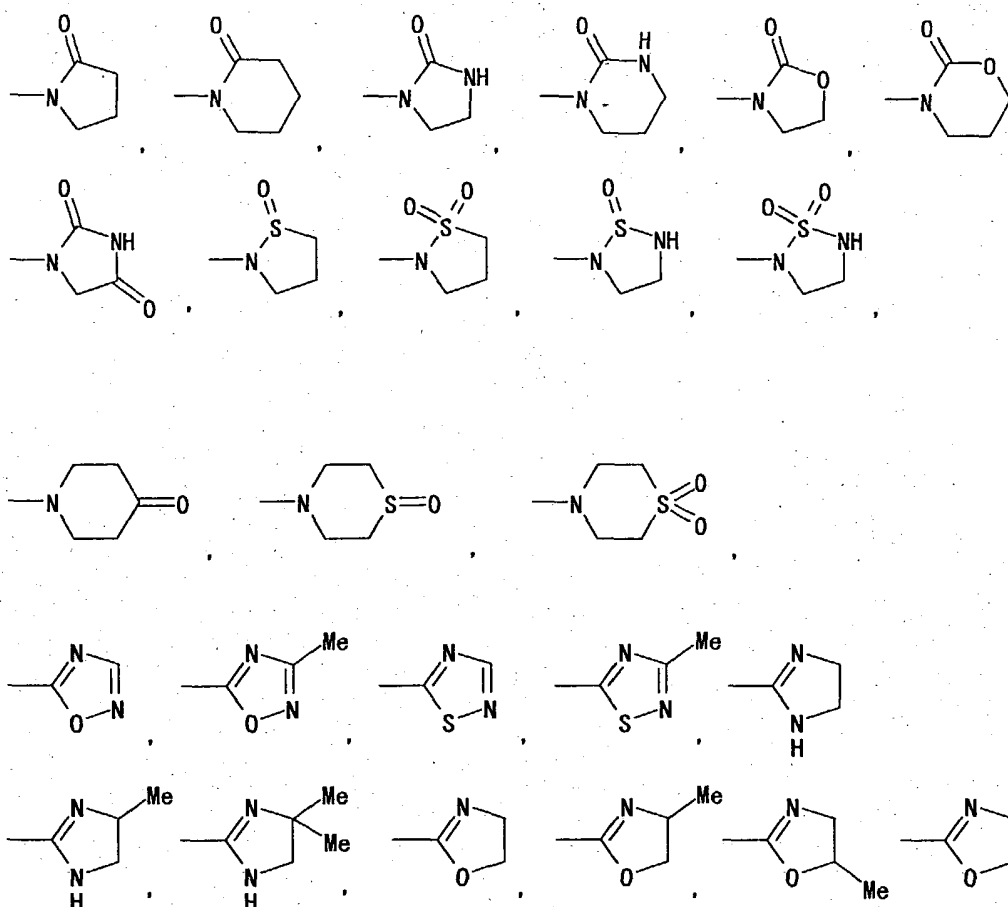
Particularly preferably, it is piperidino, 4-methylpiperidino, 2,6-dimethylpiperidino, 4-hydroxypiperidino, 1-piperazinyl, 1-pyrrolidinyl, morpholino or 4-thiomorpholinyl at  
 20 R<sup>a18</sup>, tetrahydropyranyl or 4-hydroxypiperidino at R<sup>a20</sup>, piperidino, 4-hydroxypiperidino or 3,4-dihydroxypiperidino at R<sup>a21</sup>, pyridyl or morpholino at R<sup>a24</sup>, pyridyl or 4-hydroxypiperidino at R<sup>a25</sup>, pyridyl

or thiazolyl at R<sup>a26</sup> and at R<sup>a27</sup> and R<sup>a28</sup>, it is 1-(methylsulfonyl)piperidin-4-yl, 3-hydroxypyrrolidinyl, 3-hydroxypiperidino, 4-hydroxypiperidino, 3,4-dihydroxypiperidino, 4-methoxypiperidino, 4-carboxypiperidino, 4-(hydroxymethyl)piperidino, 2-oxopiperidino, 4-oxopiperidino, 2,2,6,6-tetramethylpiperidino, 2,2,6,6-tetramethyl-4-hydroxypiperidino, 4-methylsulfonylpiperazinyl, 1-oxothiomorpholin-4-yl or 1,1-dioxothiomorpholin-4-yl, and 2-oxazolin-2-yl at R<sup>a22</sup> and R<sup>a23</sup>.

10       The heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from group D is that wherein the above-defined heterocyclic group is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted heterocyclic group. The substituent(s) is(are) selected from the substituent(s) of  
15 the above-mentioned group D (substituents shown under (a) to (q)).

Examples of the group D here include the substituent(s) exemplified for C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D.

Examples of heterocyclic group optionally substituted by 1  
20 to 5 substituent(s) selected from group D include 2-pyridyl, 3-pyridyl, 4-pyridyl, 3-fluoropyridin-4-yl, 3-chloropyridin-4-yl, 4-chloropyridin-3-yl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, 2-thienyl, 3-thienyl, furyl, oxazolyl, 2-  
25 methyloxazol-4-yl, isoxazolyl, thiazolyl, 2-methylthiazol-4-yl, 2,5-dimethylthiazol-4-yl, 2,4-dimethylthiazol-5-yl, isothiazolyl, thiadiazolyl, pyrrolinyl, pyrrolidinyl, imidazolidinyl, piperidyl, N-methylpiperidin-4-yl, N-(tert-butoxycarbonyl)piperidin-4-yl, N-acetylpiperidin-4-yl, N-methylsulfonylpiperidin-4-yl, piperazinyl,  
30 morpholinyl, thiomorpholinyl, tetrahydropyranyl, quinolyl, isoquinolyl, quinazolinyl, quinoxalyl, phthalazinyl, cinnolinyl, naphthyridinyl, 5,6,7,8-tetrahydroquinolyl, indolinyl, benzimidazolyl, indolinyl, benzofuranyl, benzothienyl, benzoxazolyl, benzothiazolyl



5 and the like.

In addition, the heterocyclic group may be substituted at the 3-, 4-, 5- or 6-position of 2-pyridyl, at the 2-, 4-, 5- or 6-position of 3-pyridyl, at the 2-, 3-, 5- or 6-position of 4-pyridinyl, at the 3-, 4- or 5-position of 2-thienyl, or at the 2-,  
 10 4- or 5-position of 3-thienyl, by fluorine atom, chlorine atom, bromine atom, nitro, methyl, tert-butyl, carboxyl, trifluoromethyl, hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl, methylsulfonyl, amino or acetyl amino.

15 At Z and Z', the heterocyclic moiety is preferably a heterocyclic group which is a 5-membered or 6-membered monocyclic group. Examples thereof include pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, 2-oxopyrrolidinyl, 2-oxopiperidyl, pyrazolyl, imidazolyl, 2-imidazolinyl, 2-oxoimidazolidinyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl,  
 20 oxazolyl, isoxazolyl, 2-oxazolynyl, thiazolyl, isothiazolyl, 1,1-dioxoisothiazolidinyl, thiadiazolyl, pyrrolidinyl, piperidyl,

piperazinyl, morpholinyl, thiomorpholinyl, tetrahydropyranyl,  $\Delta^2$ -1,2,4-oxadiazolyl, 5-oxo- $\Delta^2$ -1,2,4-oxadiazolyl, 5-oxo- $\Delta^2$ -1,2,4-thiadiazolyl and 2-oxo-3H-1,2,3,5-oxathiadiazolyl. The group D here is preferably the above-defined halogen atom, nitro, the  
 5 above-defined optionally substituted  $C_{1-6}$  alkyl,  
 $-(CH_2)_t-COOR^{a19}$ ,  $-(CH_2)_t-CONR^{a27}R^{a28}$ ,  $-(CH_2)_t-OR^{a20}$ ,  $-(CH_2)_t-NR^{a29}CO-R^{a24}$ ,  
 $-(CH_2)_t-S(O)_q-R^{a25}$  or  $-(CH_2)_t-SO_2-NHR^{a26}$ .

Examples of heterocyclic group optionally substituted by 1 to 5 substituent(s) selected from group D preferably include  
 10 piperidino, 4-hydroxypiperidino, 2-oxopiperidin-1-yl, 1-piperazinyl, 1-pyrrolidinyl, 2-oxopyrrolidin-1-yl, morpholino, 4-thiomorpholinyl, 4-tetrahydropyranyl, 3-pyridyl, 2-pyrimidinyl, 2-imidazolin-2-yl, 2-oxoimidazolidin-1-yl, 2-oxooxazolidin-1-yl, 5-tetrazolyl, 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 2-  
 15 methylthiazol-4-yl, 5-methylthiazol-2-yl, 2-aminothiazol-4-yl, 3-methyl-1,2,4-oxadiazol-5-yl, 1,1-dioxoisothiazolidin-2-yl, 4,4-dimethyl- $\Delta^2$ -oxazolin-2-yl, 2-thienyl, 5-chlorothiophen-2-yl, 5-methyloxazol-2-yl, 5-oxo- $\Delta^2$ -1,2,4-oxadiazolin-3-yl, 5-oxo- $\Delta^2$ -1,2,4-thiadiazolin-3-yl and 2-oxo-3H-1,2,3,5-oxathiazolin-4-yl.

20 Particularly preferably, it is pyridyl, pyrimidinyl, tetrazolyl, thienyl, piperidyl, 2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 2-imidazolin-2-yl, 2-oxoimidazolidin-1-yl, 2-oxooxazolidin-1-yl, 2-methylthiazol-4-yl, 5-methylthiazol-2-yl, 2-aminothiazol-4-yl, 3-methyl-1,2,4-oxadiazol-5-yl, 1,1-dioxoisothiazolidin-2-yl, 4,4-dimethyl- $\Delta^2$ -oxazolin-2-yl, 5-chlorothiophen-2-yl, 5-methyloxazol-2-yl, 5-oxo- $\Delta^2$ -1,2,4-oxadiazolin-3-yl, 5-oxo- $\Delta^2$ -1,2,4-thiadiazolin-3-yl or 2-oxo-3H-1,2,3,5-oxathiazolin-4-yl, more preferably 2-oxopyrrolidin-1-yl.

30 The  $C_{3-8}$  cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from group C is that wherein the above-defined  $C_{3-8}$  cycloalkyl is optionally substituted by the 1 to 5 substituent(s) selected from hydroxyl group, the above-defined halogen atom, the above-defined  $C_{1-6}$  alkyl and the above-defined  
 35  $C_{1-6}$  alkoxy, which may be unsubstituted. Examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, 4-fluorocyclohexyl, 2-methylcyclopentyl, 3-methylcyclohexyl, 4-methylcyclohexyl, 4,4-dimethylcyclohexyl, 3,5-dimethylcyclohexyl,

4-tert-butylcyclohexyl, 4-hydroxycyclohexyl, 4-methoxycyclohexyl and 2,3,4,5,6-pentafluorocyclohexyl.

The cycloalkyl moiety is preferably cyclopentyl or cyclohexyl, particularly preferably cyclohexyl.

5        At the ring Cy and ring Cy', the C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from group C is preferably cyclopentyl, cyclohexyl, 4-fluorocyclohexyl, 4-methylcyclohexyl, 4,4-dimethylcyclohexyl, 4-tert-butylcyclohexyl, 4-hydroxycyclohexyl or 4-methoxycyclohexyl, more preferably  
10 cyclopentyl or cyclohexyl, particularly preferably cyclohexyl.

The C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from the above group B is that wherein the above-defined C<sub>3-8</sub> cycloalkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted cycloalkyl. The  
15 substituents are selected from the above group B.

Specific examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, 4-fluorocyclohexyl, 2-methylcyclopentyl, 3-methylcyclohexyl, 4-methylcyclohexyl, 4,4-dimethylcyclohexyl, 3,5-dimethylcyclohexyl, 4-tert-  
20 butylcyclohexyl, 4-hydroxycyclohexyl, 4-methoxycyclohexyl and 2,3,4,5,6-pentafluorocyclohexyl.

Also exemplified are those wherein cyclopentyl or cyclohexyl is substituted by fluorine atom, chlorine atom, bromine atom, nitro, methyl, tert-butyl, carboxyl, trifluoromethyl, hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl, methylsulfonyl or acetylamino.  
25

At cycloalkyl moiety, it is preferably cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl. As the C<sub>3-8</sub> cycloalkyl  
30 optionally substituted by 1 to 5 substituent(s) selected from the above group B, it is particularly preferably cyclopropyl, cyclobutyl, cyclohexyl or 4-hydroxycyclohexyl at R<sup>a27</sup> and R<sup>a28</sup>.

The C<sub>3-8</sub> cycloalkyl optionally substituted by 1 to 5 substituent(s) selected from group D is that wherein the above-  
35 defined C<sub>3-8</sub> cycloalkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted cycloalkyl. The substituent(s) is(are) selected from the substituent(s) of the above-mentioned group D (substituents shown under (a) to (q)).

The group D here includes the substituents recited with regard to C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D.

Examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, 4-fluorocyclohexyl, 2-methylcyclopentyl, 3-methylcyclohexyl, 4-methylcyclohexyl, 4,4-dimethylcyclohexyl, 3,5-dimethylcyclohexyl, 4-tert-butylcyclohexyl, 4-hydroxycyclohexyl, 4-methoxycyclohexyl and 2,3,4,5,6-pentafluorocyclohexyl.

10 The group D may be, for example, cyclopentyl or cyclohexyl substituted by fluorine atom, chlorine atom, bromine atom, nitro, methyl, tert-butyl, carboxyl, trifluoromethyl, hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl, methylsulfonyl or acetylamino.

15 The cycloalkyl moiety is preferably cyclopentyl or cyclohexyl, and at Z and Z', it is particularly preferably cyclohexyl.

The optionally substituted C<sub>3-8</sub> cycloalkenyl is that wherein the above-defined C<sub>3-8</sub> cycloalkenyl is optionally substituted by substituent(s) selected from hydroxyl group, the above-defined halogen atom, the above-defined C<sub>1-6</sub> alkyl and the above-defined C<sub>1-6</sub> alkoxy, which may be unsubstituted. Examples thereof include cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclopentadienyl, cyclohexenyl, 4-fluoro-2-cyclohexenyl, 4-methyl-2-cyclohexenyl, 4-methyl-3-cyclohexenyl, 2,4-cyclohexadien-1-yl, 2,5-cyclohexadien-1-yl, cycloheptenyl and cyclooctenyl and the like, but do not include aryl (e.g., phenyl) or completely saturated cycloalkyl.

The optionally substituted C<sub>3-8</sub> cycloalkenyl is particularly preferably cyclohexenyl at the ring Cy.

The C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group B is that wherein the above-defined C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted arylalkyl. The substituent(s) is(are) selected from the above-mentioned group B.

Examples thereof include benzyl, 1-naphthylmethyl, 2-naphthylmethyl, phenethyl, 3-phenylpropyl, 2-phenylpropyl, 3-fluorobenzyl, 4-fluorobenzyl, 3-chlorobenzyl, 4-chlorobenzyl,

2,4-dichlorobenzyl, 3,5-dichlorobenzyl, pentafluorobenzyl, 4-methylbenzyl, 4-tert-butylbenzyl, 2-trifluoromethylbenzyl, 4-trifluoromethylbenzyl, 4-nitrobenzyl, 4-cyanobenzyl, 4-acetylbenzyl, 4-carboxylbenzyl, 4-carbamoylbenzyl, 4-aminobenzyl, 5 4-dimethylaminobenzyl, 4-acetylaminobenzyl, 4-(methylsulfonylamino)benzyl, 4-methoxybenzyl, 3,4,5-trimethoxybenzyl, 4-methylthiobenzyl, 4-methylsulfonylbenzyl, 4-aminosulfonylbenzyl, 3-nitro-4-methoxybenzyl and 4-nitro-3-methoxybenzyl.

10 The C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl moiety is preferably benzyl or phenethyl, particularly preferably benzyl. The group B is preferably the above-defined halogen atom, nitro, the above-defined C<sub>1-6</sub> alkyl, the above-defined halogenated C<sub>1-6</sub> alkyl or -(CH<sub>2</sub>)<sub>r</sub>-OR<sup>b1</sup>. Examples thereof include fluorine atom, chlorine 15 atom, nitro, methyl, tert-butyl, trifluoromethyl, methoxy or trifluoromethoxy, particularly preferably fluorine atom or chlorine atom.

The specific C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group B at R<sup>a12</sup> and R<sup>a13</sup> is 20 preferably benzyl, phenethyl, 3-chlorobenzyl, 4-chlorobenzyl, 4-tert-butylbenzyl or 3-trifluoromethylbenzyl, it is preferably benzyl at R<sup>a1</sup>, R<sup>a19</sup>, R<sup>a27</sup>, R<sup>a28</sup>, R<sup>a31</sup> and R<sup>b5</sup>, it is preferably benzyl, phenethyl, 4-fluorobenzyl, 2-chlorobenzyl, 3-chlorobenzyl, 4-chlorobenzyl, 4-tert-butylbenzyl or 4-trifluoromethylbenzyl at 25 R<sup>a20</sup>, and 4-chlorobenzyl, 3,5-dichlorobenzyl or 4-trifluoromethylbenzyl at R<sup>a22</sup> and R<sup>a23</sup>.

It is particularly preferably benzyl at other substituents.

The C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D is that wherein the above- 30 defined C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted aryl. The substituent(s) is(are) selected from the substituent(s) of the above-mentioned group D (substituents shown under (a) to (q)).

Examples of group D include fluorine atom, chlorine atom, 35 bromine atom, nitro, cyano, methyl, ethyl, propyl, isopropyl, tert-butyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, methoxymethyl, 2-carboxylethyl, methoxycarbonylmethyl, ethoxycarbonylmethyl, acetyl, carboxyl, methoxycarbonyl,

ethoxycarbonyl, carbamoyl, methylaminocarbonyl, isopropylaminocarbonyl, dimethylaminocarbonyl, diethylaminocarbonyl, (2-hydroxyethyl)aminocarbonyl, (carboxymethyl)aminocarbonyl, hydroxyl group, methoxy, ethoxy,  
 5 isopropoxy, hydroxymethoxy, carboxymethoxy, (dimethylaminocarbonyl)methoxy, amino, methylamino, dimethylamino, diethylamino, acetylamino, methylsulfonylamino, methylthio, methylsulfonyl, methylsulfinyl, aminosulfonyl, methylaminosulfonyl and dimethylaminosulfonyl.

10 Examples of C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D include benzyl, 1-naphthylmethyl, 2-naphthylmethyl, phenethyl, 3-phenylpropyl, 2-phenylpropyl, 3-fluorobenzyl, 4-fluorobenzyl, 3-chlorobenzyl, 4-chlorobenzyl, 2,4-dichlorobenzyl, 3,5-dichlorobenzyl, 4-bromobenzyl, 4-nitrobenzyl, pentafluorobenzyl, 4-methylbenzyl, 4-tert-butylbenzyl, 2-trifluoromethylbenzyl, 4-trifluoromethylbenzyl, 4-(hydroxymethyl)benzyl, 4-(methoxymethyl)benzyl, 4-(2-carboxylethyl)benzyl, 3-carboxylbenzyl, 4-carboxylbenzyl, 4-methoxybenzyl, 3,4,5-trimethoxybenzyl, 4-carbamoylbenzyl, 4-methylthiobenzyl, 4-(dimethylaminocarbonyl)benzyl, 4-methylsulfonylbenzyl, 4-(acetylamino)benzyl, 4-cyanobenzyl, 4-acetylbenzyl, 4-aminobenzyl, 4-dimethylaminobenzyl, 4-(methylsulfonylamino)benzyl, 4-methylsulfinylbenzyl, 4-aminosulfonylbenzyl, (3-nitro-4-methoxyphenyl)methyl and (4-nitro-3-methoxyphenyl)methyl.  
 25

At Z and Z', the C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl moiety is preferably benzyl or phenethyl, and the group D here is preferably the above-defined halogen atom, nitro, the above-defined optionally substituted C<sub>1-6</sub> alkyl,  $-(CH_2)_t-COOR^{a19}$ ,  $-(CH_2)_t-CONR^{a27}R^{a28}$ ,  
 30  $-(CH_2)_t-OR^{a20}$ ,  $-(CH_2)_t-NR^{a29}CO-R^{a24}$ ,  $-(CH_2)_t-S(O)_q-R^{a25}$  or  $-(CH_2)_t-SO_2-NHR^{a26}$ .

The C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D is preferably benzyl, 3-fluorobenzyl, 4-fluorobenzyl, 3-chlorobenzyl, 4-chlorobenzyl, 3,5-dichlorobenzyl, 4-bromobenzyl, 4-nitrobenzyl, 4-methylbenzyl, 4-tert-butylbenzyl, 2-trifluoromethylbenzyl, 4-trifluoromethylbenzyl, 4-(hydroxymethyl)benzyl, 4-(methoxymethyl)benzyl, 4-(2-carboxylethyl)benzyl, 3-



carboxylbenzyl, 4-carboxylbenzyl, 4-methoxybenzyl, 3,4,5-trimethoxybenzyl, 4-carbamoylbenzyl, 4-methylthiobenzy, 4-(dimethylaminocarbonyl)benzyl, 4-methylsulfonylbenzyl, 4-acetylaminobenzy, 4-methylsulfinylbenzyl or 4-aminosulfonylbenzyl.

It is particularly preferably the above-defined halogen atom, the above-defined optionally substituted  $C_{1-6}$  alkyl,  $-(CH_2)_t-COOR^{a19}$ ,  $-(CH_2)_t-CONR^{a27}R^{a28}$ ,  $-(CH_2)_t-OR^{a20}$  or  $-(CH_2)_t-S(O)_q-R^{a25}$ . Examples thereof include fluorine atom, chlorine atom, bromine atom, nitro, methyl, tert-butyl, carboxyl, trifluoromethyl, hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl, methylsulfonyl and acetyl amino. It is more preferably fluorine atom, chlorine atom, methyl, tert-butyl, carboxyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl or methylsulfonyl, most preferably fluorine atom or chlorine atom.

The heterocycle  $C_{1-6}$  alkyl optionally substituted by 1 to 5 substituent(s) selected from group B is that wherein the above-defined heterocycle  $C_{1-6}$  alkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted heterocycle  $C_{1-6}$  alkyl. The substituent(s) is(are) selected from the above-mentioned group B.

Examples thereof include 2-pyridylmethyl, 3-pyridylmethyl, 2-chloropyridin-4-ylmethyl, 4-pyridylmethyl, pyrrolylmethyl, imidazolylmethyl, 2-thienylmethyl, 3-thienylmethyl, 2-furylmethyl, 2-oxazolylmethyl, 5-isothiazolylmethyl, 2-methyloxazol-4-ylmethyl, 2-thiazolylmethyl, 4-thiazolylmethyl, 5-thiazolylmethyl, 2-methylthiazol-4-ylmethyl, 2-methylthiazol-5-ylmethyl, 2,5-dimethylthiazol-4-ylmethyl, 4-methylthiazol-2-ylmethyl, 2,4-dimethylthiazol-5-ylmethyl, 2-isothiazolylmethyl, 2-pyrrolinylmethyl, pyrrolidinylmethyl, piperidylmethyl, 4-piperidylmethyl, 1-methylpiperidin-4-ylmethyl, 4-hydroxypiperidinomethyl, 3-hydroxypyrrolidinylmethyl, 2-(4-hydroxypiperidino)ethyl, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyl, 1-acetylpiperidin-4-ylmethyl, 1-methylsulfonylpiperidin-4-ylmethyl, piperazinylmethyl, morpholinomethyl, thiomorpholinylmethyl, 1-

tetrahydropyranylmethyl, 2-quinolylmethyl, 1-isoquinolylmethyl and the like.

The heterocyclic moiety is preferably a heterocyclic group which is a 5-membered or 6-membered monocyclic group. Examples thereof include pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyrrolidinyl, piperidyl, piperazinyl, morpholinyl, thiomorpholinyl and tetrahydropyranyl, and the alkyl moiety thereof is preferably straight chain alkyl having 1 to 4 carbon atoms. The group B here is preferably the above-defined halogen atom, the above-defined C<sub>1-6</sub> alkyl, the above-defined halogenated C<sub>1-6</sub> alkyl, the above-defined C<sub>1-6</sub> alkanoyl,  $-(CH_2)_r-COOR^{b1}$ ,  $-(CH_2)_r-CONR^{b1}R^{b2}$  or  $-(CH_2)_r-OR^{b1}$ .

Examples of heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group B preferably include 2-pyridylmethyl, 3-pyridylmethyl, 2-chloropyridin-4-ylmethyl, 4-pyridylmethyl, piperidin-4-ylmethyl, 1-methylpiperidin-4-ylmethyl, 2-(4-hydroxypiperidino)ethyl, 1-acetylpiperidin-4-ylmethyl, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyl, 1-(methylsulfonyl)piperidin-4-ylmethyl, 2-thiazolylmethyl, 4-thiazolylmethyl, 2-methylthiazolin-4-ylmethyl, 2,4-dimethylthiazolin-5-ylmethyl and 4-methylthiazol-2-ylmethyl. Particularly preferably, it is 2-pyridylmethyl, 3-pyridylmethyl, 2-chloropyridin-4-ylmethyl, 4-pyridylmethyl, piperidin-4-ylmethyl, 1-methylpiperidin-4-ylmethyl, 2-(4-hydroxypiperidino)ethyl, 1-acetylpiperidin-4-ylmethyl, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyl, 1-(methylsulfonyl)piperidin-4-ylmethyl, 2-methylthiazolin-4-ylmethyl, 2,4-dimethylthiazolin-5-ylmethyl or 4-methylthiazol-2-ylmethyl at R<sup>a20</sup>, 2-pyridylmethyl at R<sup>a22</sup> and R<sup>a23</sup>, and 4-pyridylmethyl or 4-methylthiazol-2-ylmethyl at R<sup>a27</sup> and R<sup>a28</sup>.

The heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D is that wherein the above-defined heterocycle C<sub>1-6</sub> alkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted heterocycle C<sub>1-6</sub> alkyl. The substituent(s) is(are) selected from the above-mentioned group D (substituents shown under (a) to (q)).

Examples of group D here include fluorine atom, chlorine atom, bromine atom, nitro, cyano, methyl, ethyl, propyl, isopropyl, tert-butyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, methoxymethyl, 2-carboxylethyl, methoxycarbonylmethyl, ethoxycarbonylmethyl, acetyl, carboxyl, methoxycarbonyl, ethoxycarbonyl, carbamoyl, methylaminocarbonyl, isopropylaminocarbonyl, dimethylaminocarbonyl, diethylaminocarbonyl, (2-hydroxyethyl)aminocarbonyl, (carboxylmethyl)aminocarbonyl, hydroxyl group, methoxy, ethoxy, isopropoxy, hydroxymethoxy, carboxylmethoxy, (dimethylaminocarbonyl)methoxy, amino, methylamino, dimethylamino, diethylamino, acetylamino, methylsulfonylamino, methylthio, methylsulfonyl, methylsulfinyl, aminosulfonyl, methylaminosulfonyl and dimethylaminosulfonyl.

Examples of heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D include 2-pyridylmethyl, 3-pyridylmethyl, 2-chloropyridin-4-ylmethyl, 4-pyridylmethyl, pyrrolylmethyl, imidazolylmethyl, 2-thienylmethyl, 3-thienylmethyl, 2-furylmethyl, 2-oxazolylmethyl, 5-isothiazolylmethyl, 2-methyloxazol-4-ylmethyl, 2-thiazolylmethyl, 4-thiazolylmethyl, 5-thiazolylmethyl, 2-methylthiazol-4-ylmethyl, 2-methylthiazol-5-ylmethyl, 2,5-dimethylthiazol-4-ylmethyl, 4-methylthiazol-2-ylmethyl, 2,4-dimethylthiazol-5-ylmethyl, 2-isothiazolylmethyl, 2-pyrrolinylmethyl, pyrrolidinylmethyl, piperidylmethyl, 4-piperidylmethyl, 1-methylpiperidin-4-ylmethyl, 4-hydroxypiperidinomethyl, 2-(4-hydroxypiperidino)ethyl, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyl, 1-acetylpiperidin-4-ylmethyl, 1-methylsulfonylpiperidin-4-ylmethyl, piperazinylmethyl, morpholinomethyl, thiomorpholinylmethyl, 1-tetrahydropyranylmethyl, 2-quinolylmethyl, 1-isoquinolylmethyl, and the like.

Preferable heterocyclic moiety at Z and Z' is heterocyclic group which is 5-membered or 6-membered monocyclic group. Examples of the heterocyclic moiety include pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isooxazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyrrolidinyl, piperidyl, piperazinyl, morpholinyl, thiomorpholinyl and

tetrahydropyranyl, and the alkyl moiety is preferably straight chain alkyl having 1 to 4 carbon atoms, particularly methyl (i.e., methylene).

Preferable group D is the above-defined halogen atom, 5 nitro, the above-defined optionally substituted C<sub>1-6</sub> alkyl, - (CH<sub>2</sub>)<sub>t</sub>-COOR<sup>a19</sup>, - (CH<sub>2</sub>)<sub>t</sub>-CONR<sup>a27</sup>R<sup>a28</sup>, - (CH<sub>2</sub>)<sub>t</sub>-OR<sup>a20</sup>, - (CH<sub>2</sub>)<sub>t</sub>-NR<sup>a29</sup>CO-R<sup>a24</sup>, - (CH<sub>2</sub>)<sub>t</sub>-S(O)<sub>q</sub>-R<sup>a25</sup> or - (CH<sub>2</sub>)<sub>t</sub>-SO<sub>2</sub>-NHR<sup>a26</sup>.

Preferable examples of heterocycle C<sub>1-6</sub> alkyl optionally substituted by 1 to 5 substituent(s) selected from group D 10 include 2-pyridylmethyl, 3-pyridylmethyl, 2-chloropyridin-4-ylmethyl, 4-pyridylmethyl, piperidin-4-ylmethyl, 1-methylpiperidin-4-ylmethyl, 4-hydroxypiperidinomethyl, 2-(4-hydroxypiperidino)ethyl, 1-acetylpiperidin-4-ylmethyl, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyl, 1-(methylsulfonyl)piperidin- 15 4-ylmethyl, 2-thiazolylmethyl, 4-thiazolylmethyl, 2-methylthiazolin-4-ylmethyl, 2,4-dimethylthiazolin-5-ylmethyl and 4-methylthiazol-2-ylmethyl.

Particularly preferred is 4-hydroxypiperidinomethyl.

The C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl optionally substituted by 1 20 to 5 substituent(s) selected from the above group B is that wherein the above-defined C<sub>3-8</sub> cycloalkyl C<sub>1-6</sub> alkyl is optionally substituted by 1 to 5 substituent(s), and includes unsubstituted cycloalkylalkyl. The substituents are selected from the above group B.

25 Specific examples thereof include cyclopropylmethyl, cyclobutylmethyl, cyclopentylmethyl, cyclohexylmethyl, 2-(cyclopentyl)ethyl, 2-(cyclohexyl)ethyl, cycloheptylmethyl, 4-fluorocyclohexylmethyl, 2-methylcyclopentylmethyl, 3-methylcyclohexylmethyl, 4-methylcyclohexylmethyl, 4,4- 30 dimethylcyclohexylmethyl, 3,5-dimethylcyclohexylmethyl, 4-tert-butylcyclohexylmethyl, 4-hydroxycyclohexylmethyl, 4-methoxycyclohexylmethyl and 2,3,4,5,6-pentafluorocyclohexylmethyl.

Also exemplified are those wherein cyclopentylmethyl or cyclohexylmethyl is substituted by fluorine atom, chlorine atom, 35 bromine atom, nito, methyl, tert-butyl, carboxyl, trifluoromethyl, hydroxymethyl, methoxymethyl, 2-carboxylethyl, methoxy, carbamoyl, methylthio, dimethylaminocarbonyl, methylsulfonyl or acetylamino.

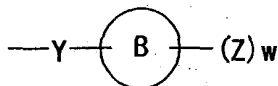
At cycloalkyl moiety, it is preferably cyclopentylmethyl

or cyclohexylmethyl, and at  $R^{a20}$ ,  $R^{a27}$  and  $R^{a28}$ , it is particularly preferably cyclohexylmethyl.

The carboxyl-protecting group only needs to be suitable for reaction conditions, and is capable of protecting and  
5 deprotecting and may be, for example, methyl; substituted methyl group such as methoxymethyl, methylthiomethyl, 2-tetrahydropyranyl, methoxyethoxymethyl, benzyloxymethyl, phenacyl, diacylmethyl, phthalimidomethyl etc.; ethyl; substituted ethyl group such as 2,2,2-trichloroethyl, 2-chloroethyl, 2-  
10 (trimethylsilyl)ethyl, 2-methylthioethyl, 2-(p-toluenesulfonyl)ethyl, t-butyl etc.; benzyl; substituted benzyl group such as diphenylmethyl, triphenylmethyl, p-nitrobenzyl, 4-picolyl, p-methoxybenzyl, 2-(9,10-dioxo)anthrylmethyl etc.; silyl group such as trimethylsilyl, t-butyldimethylsilyl,  
15 phenyldimethylsilyl etc.; and the like.

Preferred are industrially effective protecting groups and specifically preferred as  $R^{a36}$  are methyl and ethyl.

In formula [I], X is preferably

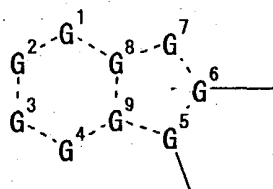


wherein each symbol is as defined above.

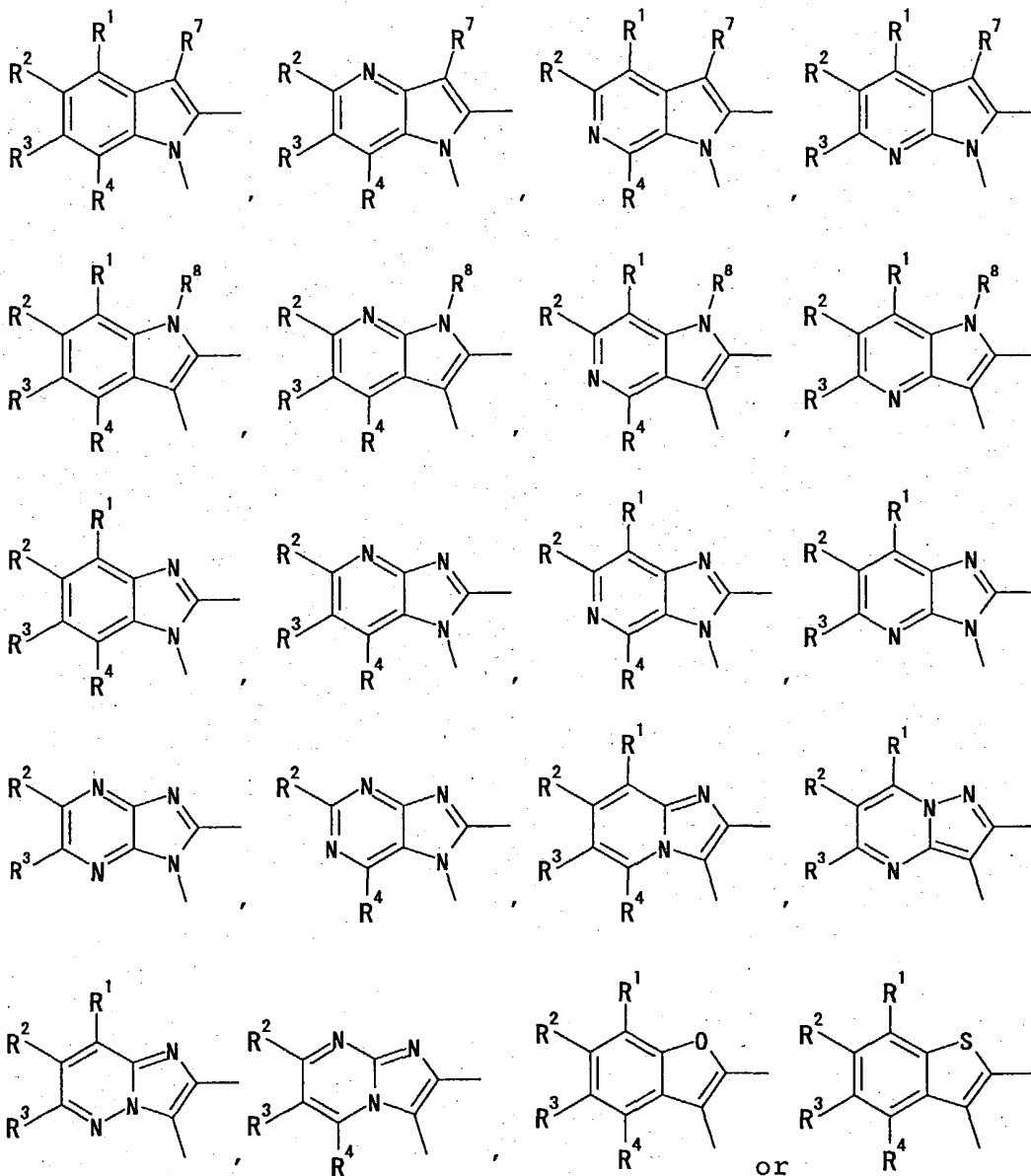
$G^1$ ,  $G^2$ ,  $G^3$  and  $G^4$  are each preferably  $(C-R^1)$ ,  $(C-R^2)$ ,  $(C-R^3)$  and  $(C-R^4)$ ,  $G^5$  is preferably a nitrogen atom, and  $G^6$ ,  $G^8$  and  $G^9$  are  
25 preferably a carbon atom.  $G^7$  is preferably  $C(-R^7)$  or unsubstituted nitrogen atom, wherein  $R^7$  is preferably hydrogen atom.

A preferable combination is  $G^2$  of  $(C-R^2)$  and  $G^6$  of a carbon atom, particularly preferably  $G^2$  of  $(C-R^2)$ ,  $G^6$  of a carbon atom  
30 and  $G^5$  of a nitrogen atom, most preferably  $G^2$  of  $(C-R^2)$ ,  $G^6$  of a carbon atom,  $G^5$  of a nitrogen atom and  $G^7$  of unsubstituted nitrogen atom.

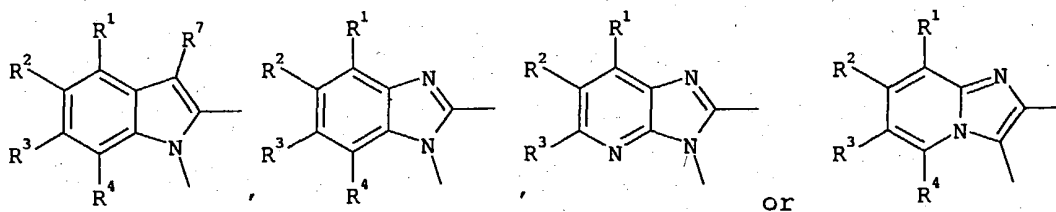
In formulas [I] and [II], 1 to 4 of  $G^1$  to  $G^9$  in the moiety



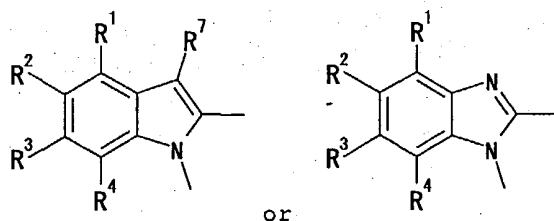
is(are) preferably a nitrogen atom, specifically preferably



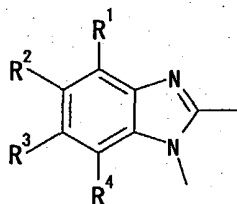
particularly preferably



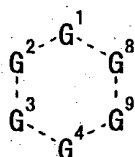
more preferably



most preferably



5 It is also a preferable embodiment wherein the



moiety is aromatic ring.

$R^1$  and  $R^4$  are preferably hydrogen atom.  $R^2$  is preferably carboxyl,  $-\text{COOR}^{a1}$ ,  $-\text{CONR}^{a2}\text{R}^{a3}$ ,  $-\text{SO}_2\text{R}^{a7}$  (each symbol is as defined above) or heterocyclic group having 1 to 4 heteroatom(s) selected  
10 from an oxygen atom, a nitrogen atom and a sulfur atom, particularly preferably carboxyl,  $-\text{COOR}^{a1}$  or  $-\text{SO}_2\text{R}^{a7}$ , more preferably carboxyl or  $-\text{COOR}^{a1}$ , most preferably carboxyl.  $R^3$  is preferably hydrogen atom or  $-\text{OR}^{a6}$  ( $\text{R}^{a6}$  is as defined above), particularly preferably hydrogen atom.

15  $\text{R}^{a1}$  is preferably optionally substituted  $\text{C}_{1-6}$  alkyl.

When  $\text{R}^2$  is carboxyl or  $-\text{COOR}^{a1}$ , at least one of  $\text{R}^1$ ,  $\text{R}^3$  and  $\text{R}^4$  is preferably hydroxyl group, halogen atom (particularly fluorine atom, chlorine atom) or  $-\text{OR}^{a6}$  (wherein  $\text{R}^{a6}$  is preferably hydrogen atom or methyl).

20 The ring Cy and ring Cy' are preferably cyclopentyl, cyclohexyl, cycloheptyl, tetrahydrothiopyranyl or piperidino, particularly preferably cyclopentyl, cyclohexyl or cycloheptyl, more preferably cyclohexyl.

The ring A and ring A' are preferably phenyl, pyridyl,  
25 pyrazinyl, pyrimidinyl, pyridazinyl, cyclohexyl, cyclohexenyl, furyl or thienyl, particularly preferably phenyl, pyridyl,

pyrazinyl, pyrimidinyl or pyridazinyl, more preferably phenyl or pyridyl, and most preferably phenyl.

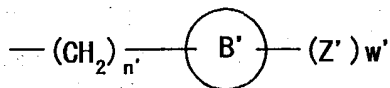
The ring B and ring B' are preferably C<sub>1-6</sub> aryl or heterocyclic group, specifically preferably, phenyl, pyridyl,  
 5 pyrazinyl, pyrimidinyl, pyridazinyl, 1,3,5-triazinyl, pyrrolyl, pyrazolyl, imidazolyl, 1,2,4-triazolyl, tetrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl or thiadiazolyl, particularly preferably phenyl, pyridyl,  
 pyrimidinyl, 1,3,5-triazinyl or thiazolyl, more preferably,  
 10 phenyl, pyridyl or thiazolyl, and most preferably phenyl or thiazolyl.

With regard to R<sup>5</sup> and R<sup>6</sup>, one of them is preferably hydrogen atom and the other is halogen atom, particularly fluorine atom. Alternatively, the both are preferably hydrogen  
 15 atoms. When ring A is phenyl, R<sup>5</sup> and R<sup>6</sup> preferably are present at an ortho position from G<sup>6</sup>. The same applies to R<sup>5'</sup> and R<sup>6'</sup>.

Y is preferably -(CH<sub>2</sub>)<sub>m</sub>-O-(CH<sub>2</sub>)<sub>n</sub>-, -NHCO<sub>2</sub>-, -CONH-CHR<sup>a14</sup>-,  
 -(CH<sub>2</sub>)<sub>m</sub>-NR<sup>a12</sup>-(CH<sub>2</sub>)<sub>n</sub>-, -CONR<sup>a13</sup>-(CH<sub>2</sub>)<sub>n</sub>-, -O-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>- or  
 -(CH<sub>2</sub>)<sub>n</sub>-NR<sup>a12</sup>-CHR<sup>a15</sup>- (each symbol is as defined above), more  
 20 preferably, -(CH<sub>2</sub>)<sub>m</sub>-O-(CH<sub>2</sub>)<sub>n</sub>- or -O-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>-, most preferably -O-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>-.

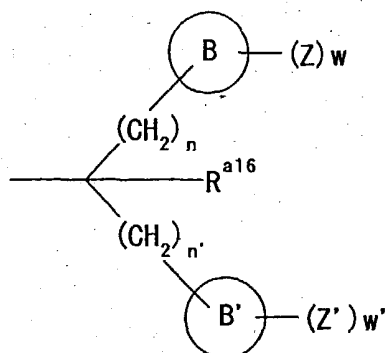
The l, m and n are preferably 0 or an integer of 1 to 4, particularly preferably 0, 1 or 2, at Y. In -(CH<sub>2</sub>)<sub>m</sub>-O-(CH<sub>2</sub>)<sub>n</sub>-,  
 m=n=0 or m=0 and n=1 is more preferable, most preferably m=n=0.  
 25 In -O-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>-, m=n=0, m=0 and n=1, m=1 and n=0 or m=1 and n=1 is more preferable, most preferably m=n=0.

When Y is -O-(CH<sub>2</sub>)<sub>m</sub>-CR<sup>a15</sup>R<sup>a16</sup>-(CH<sub>2</sub>)<sub>n</sub>-, R<sup>a16</sup> is preferably hydrogen atom, R<sup>a15</sup> is preferably



30 wherein the





moiety is preferably symmetric. The preferable mode of  $n$ , ring B, Z and  $w$  and the preferable mode of  $n'$ , ring B', Z' and  $w'$  are the same.

5 When ring A is phenyl, X or Y is preferably present at the para-position relative to  $G^6$ . When ring B and ring B' are phenyl, Z is preferably present at the ortho or meta-position relative to Y. It is preferable that the 3-position on phenyl have one substituent or the 2-position and the 5-position on phenyl each  
10 have one substituent.

When ring B is thiazolyl, Y is preferably substituted at the 5-position, and Z is preferably substituted at the 2-position, the 4-position or the 2-position and the 4-position. Similarly, when ring B' is thiazolyl,  $(CH_2)_{n'}$  is also preferably substituted  
15 at the 5-position, and Z' is preferably substituted at the 2-position, the 4-position or the 2-position and the 4-position.

Z and Z' are preferably group D, "C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D" or "heterocyclic group optionally substituted by 1 to 5  
20 substituent(s) selected from group D", particularly preferably group D or "C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D".

More preferably, they are the above-defined halogen atom, nitro, the above-defined optionally substituted C<sub>1-6</sub> alkyl,  
25  $-(CH_2)_t-COOR^{a19}$ ,  $-(CH_2)_t-CONR^{a27}R^{a28}$ ,  $-(CH_2)_t-OR^{a20}$ ,  $-(CH_2)_t-NR^{a29}CO-R^{a24}$ ,  $-(CH_2)_t-S(O)_q-R^{a25}$  or  $-(CH_2)_t-SO_2-NHR^{a26}$ , or C<sub>6-14</sub> aryl or heterocyclic group optionally substituted by these.

With regard to Z and Z', the preferable mode of group D that directly substitutes each ring B and ring B' and the  
30 preferable mode of group D that substitutes C<sub>6-14</sub> aryl, C<sub>3-8</sub> cycloalkyl, C<sub>6-14</sub> aryl C<sub>1-6</sub> alkyl or heterocyclic group are the

same, wherein they may be the same with or different from each other.

Specific examples of the substituent preferably include fluorine atom, chlorine atom, bromine atom, nitro, cyano, methyl, ethyl, propyl, isopropyl, tert-butyl, trifluoromethyl, hydroxymethyl, 2-hydroxyethyl, methoxymethyl, 2-carboxylethyl, methoxycarbonylmethyl, ethoxycarbonylmethyl, carbamoylmethoxymethyl, (dimethylaminocarbonyl)methoxymethyl, acetyl, isovaleryl, carboxyl, methoxycarbonyl, ethoxycarbonyl, carbamoyl, methylaminocarbonyl, hydroxyaminocarbonyl, ethylaminocarbonyl, propylaminocarbonyl, isopropylaminocarbonyl, butylaminocarbonyl, isobutylaminocarbonyl, tert-butylaminocarbonyl, (4-hydroxybutyl)aminocarbonyl, (1-hydroxypropan-2-yl)aminocarbonyl, (2,3-dihydroxypropyl)-aminocarbonyl, (1,3-dihydroxypropan-2-yl)aminocarbonyl, methoxyaminocarbonyl, {2-[2-(methoxy)ethoxy]ethyl}aminocarbonyl, N-ethyl-N-methylaminocarbonyl, N-methyl-N-propylaminocarbonyl, N-isopropyl-N-methylaminocarbonyl, dimethylaminocarbonyl, diethylaminocarbonyl, (2-hydroxyethyl)aminocarbonyl, (2-hydroxy-2-methylpropan-2-yl)aminocarbonyl, (carboxylmethyl)aminocarbonyl, hydroxyl group, methoxy, ethoxy, propyloxy, isopropyloxy, butyloxy, isopentyloxy, 2-isopentenyloxy, 3-isohexenyloxy, 4-methyl-3-pentenyloxy, 2-propynyloxy, trifluoromethyloxy, hydroxymethyloxy, carboxylmethyloxy, (dimethylaminocarbonyl)-methyloxy, amino, methylamino, dimethylamino, diethylamino, acetylamino, N-acetyl-N-methylamino, ureido, isopropylcarbonylamino, isobutylcarbonylamino, tert-butylcarbonylamino, (ethylamino)carbonylamino, (isopropylamino)-carbonylamino, (dimethylamino)carbonylamino, (4-hydroxypiperidino)carbonylamino, [(4-hydroxypiperidino)methyl]-carbonylamino, [(3-hydroxypyrrolidinyl)methyl]carbonylamino, methylsulfonylamino, isopropylsulfonylamino, N-(isopropylsulfonyl)-N-methylamino, methylthio, methylsulfonyl, isopropylsulfonyl, isobutylsulfonyl, methylsulfinyl, isopropylsulfinyl, aminosulfonyl, methylaminosulfonyl, dimethylaminosulfonyl, isopropylaminosulfonyl, tert-butylaminosulfonyl, hydroxyamidino, phenyl, 3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 2,4-difluorophenyl, 3,4-

difluorophenyl, 3,4-dichlorophenyl, 3,5-dichlorophenyl, 4-chloro-  
 3-fluorophenyl, 4-chloro-2-fluorophenyl, 4-bromophenyl, 4-  
 nitrophenyl, 4-cyanophenyl, 4-methylphenyl, 4-ethylphenyl, 4-  
 propylphenyl, 4-isopropylphenyl, 4-tert-butylphenyl, 2-  
 5 trifluoromethylphenyl, 4-trifluoromethylphenyl, 4-  
 (hydroxymethyl)phenyl, 4-(2-hydroxyethyl)phenyl, 4-  
 (methoxymethyl)phenyl, 4-(2-carboxylethyl)phenyl, 4-  
 (methoxycarbonylmethyl)phenyl, 4-(ethoxycarbonylmethyl)phenyl, 4-  
 acetylphenyl, 3-carboxylphenyl, 4-carboxylphenyl, 4-  
 10 (methoxycarbonyl)phenyl, 4-(ethoxycarbonyl)phenyl, 4-  
 carbamoylphenyl, 4-(methylaminocarbonyl)phenyl, 4-  
 (isopropylaminocarbonyl)phenyl, 4-(dimethylaminocarbonyl)phenyl,  
 4-(diethylaminocarbonyl)phenyl, 4-[(2-hydroxyethyl)-  
 aminocarbonyl]phenyl, 4-[(carboxylmethyl)aminocarbonyl]phenyl, 4-  
 15 hydroxyphenyl, 4-methoxyphenyl, 3,4,5-trimethoxyphenyl, 4-  
 ethoxyphenyl, 4-propyloxyphenyl, 4-isopropyloxyphenyl, 4-  
 butyloxyphenyl, 4-isopentyloxyphenyl, 4-(2-isopentenyl)oxyphenyl,  
 4-(3-isohexenyl)oxyphenyl, 4-(4-methyl-3-pentenyl)oxyphenyl, 4-  
 (2-propenyl)oxyphenyl, 4-(trifluoromethyl)oxyphenyl, 4-  
 20 (hydroxymethyl)oxyphenyl, 4-(carboxylmethyl)oxyphenyl, 4-  
 [(dimethylaminocarbonyl)methyl]oxyphenyl, 4-aminophenyl, 4-  
 (methylamino)phenyl, 4-(dimethylaminophenyl), 4-(diethylamino)-  
 phenyl, 4-(acetylaminophenyl), 4-(methylsulfonylamino)phenyl, 4-  
 (methylthio)phenyl, 4-(methylsulfonyl)phenyl, 4-  
 25 (methylsulfinyl)phenyl, 4-(aminosulfonyl)phenyl, 4-  
 (methylaminosulfonyl)phenyl, 4-(dimethylaminosulfonyl)phenyl, 4-  
 (tert-butylaminosulfonyl)phenyl, tetrazol-5-ylphenyl, cyclohexyl,  
 benzyl, 4-chlorobenzyl, phenethyl, benzyloxy, 4-fluorobenzyloxy,  
 2-chlorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-tert-  
 30 butylbenzyloxy, 4-trifluoromethylbenzyloxy, phenethyl, 2-  
 thienyl, 2-thiazolyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 6-  
 fluoropyridin-3-yl, 5-fluoropyridin-2-yl, 6-chloropyridin-3-yl,  
 6-methylpyridin-3-yl, 2-pyrimidinyl, 5-tetrazolyl, piperidino, 2-  
 oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 2-imidazolin-2-yl, 2-  
 35 oxoimidazolidin-1-yl, 2-oxooxazolidin-1-yl, 2-methylthiazol-4-yl,  
 5-methylthiazol-2-yl, 2-aminothiazol-4-yl, 3-methyl-1,2,4-  
 oxadiazol-5-yl, 1,1-dioxoisothiazolidin-2-yl, 4,4-dimethyl- $\Delta^2$ -  
 oxazolin-2-yl, 5-chlorothiophen-2-yl, 5-methyloxazol-2-yl, 5-oxo-

$\Delta^2$ -1,2,4-oxadiazolin-3-yl, 5-oxo- $\Delta^2$ -1,2,4-thiadiazolin-3-yl, 2-oxo-3H-1,2,3,5-oxathiadiazolin-4-yl, 4-hydroxypiperidinomethyl, piperidinocarbonyl, 4-hydroxypiperidinocarbonyl, 3,4-dihydroxypiperidinocarbonyl, 1-piperazinylcarbonyl, 1-  
5 pyrrolidinylcarbonyl, morpholinocarbonyl, 4-thiomorpholinylcarbonyl, phenoxy, 2,4-dichlorophenoxy, tetrahydropyranyloxy, 2-pyridylmethyloxy, 3-pyridylmethyloxy, 2-chloropyridin-4-ylmethyloxy, 4-pyridylmethyloxy, 2-piperidylmethyloxy, 3-piperidylmethyloxy, 4-piperidylmethyloxy,  
10 1-methylpiperidin-4-ylmethyloxy, 1-acetylpiperidin-4-ylmethyloxy, 1-(tert-butoxycarbonyl)piperidin-4-ylmethyloxy, 1-(methylsulfonyl)piperidin-4-ylmethyloxy, 2-methylthiazolin-4-yloxy, 2,4-dimethylthiazolin-5-yloxy, dimethylaminocarbonylmethyloxy, piperidinocarbonylmethyloxy, 4-hydroxypiperidinocarbonylmethyloxy, 2-methylthiazol-4-yl, (2-methylthiazol-4-yl)methyloxy, (2,4-dimethylthiazol-5-yl)methyloxy, benzoyl, 3-fluorobenzoyl, 4-chlorobenzylamino, 3,5-dichlorobenzylamino, 4-trifluoromethylbenzylamino, 2-pyridylmethylamino, benzoylamino, 4-chlorobenzoylamino, 4-trifluoromethylbenzoylamino, 3,5-  
20 dichlorobenzoylamino, 3-nitro-4-methoxybenzoylamino, 4-nitro-3-methoxybenzoylamino, 3-pyridylcarbonylamino, morpholinocarbonylamino, 2-oxazolinylamino, 4-hydroxypiperidinosulfonyl, 4-methylphenylsulfonylamino, 2-thiazolylaminosulfonyl, 2-pyridylaminosulfonyl, benzylaminocarbonyl, N-benzyl-N-methylaminocarbonyl, (4-pyridylmethyl)aminocarbonyl or  
25 (cyclohexylmethyl)aminocarbonyl, 2-hydroxyethyloxy, 3-hydroxypropyloxy, 2-methoxyethoxy, 2-(2-methoxyethoxy)ethoxy, azetidylcarbonyl, 3-hydroxypyrrolidinylcarbonyl, 3-hydroxypiperidinocarbonyl, 4-hydroxypiperidinocarbonyl, 3,4-dihydroxypiperidinocarbonyl, 4-methoxypiperidinocarbonyl, 4-carboxypiperidinocarbonyl, 4-(hydroxymethyl)piperidinocarbonyl, 2-oxopiperidinocarbonyl, 4-oxopiperidinocarbonyl, 2,6-dimethylpiperidinocarbonyl, 2,2,6,6-tetramethylpiperidinocarbonyl, 2,2,6,6-tetramethyl-4-hydroxypiperidinocarbonyl, 1-  
30 oxothiomorpholin-4-ylcarbonyl, 1,1-dioxothiomorpholin-4-ylcarbonyl, 1-(methylsulfonyl)piperidin-4-ylaminocarbonyl, 4-methylsulfonylpiperazinylcarbonyl, 4-methylpiperazinylcarbonyl, N,N-bis(2-hydroxyethyl)aminocarbonyl, phenylaminocarbonyl,

cyclopropylaminocarbonyl, cyclobutylaminocarbonyl, cyclohexylaminocarbonyl, 4-hydroxycyclohexylaminocarbonyl, 4-methylthiazol-2-ylmethylaminocarbonyl, 2-(4-hydroxypiperidino)-ethyloxy, 2-pyridylmethylaminocarbonyl, 3-pyridylmethylamino-  
5 carbonyl, N-methyl-N-(4-pyridylmethyl)aminocarbonyl, cyclohexylmethyloxy, 4-hydroxypiperidinocarbonylmethyloxy and 4-methylthiazol-2-ylmethyloxy.

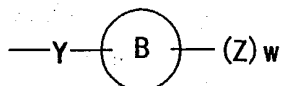
Particularly preferable examples of the substituent include fluorine atom, chlorine atom, bromine atom, nitro, cyano,  
10 methyl, hydroxymethyl, carboxyl, carbamoyl, methylaminocarbonyl, isopropylaminocarbonyl, dimethylaminocarbonyl, diethylamino-carbonyl, (2-hydroxyethyl)aminocarbonyl, (carboxymethyl)-aminocarbonyl, methoxy, 2-isopentenyl, 2-propynyl, methylthio, methylamino, dimethylamino, acetylamino,  
15 methylsulfonylamino, methylsulfonyl, aminosulfonyl, dimethylaminosulfonyl, tert-butylaminosulfonyl, phenyl, 3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 3,5-dichlorophenyl, 4-nitrophenyl, 4-methylphenyl, 4-tert-butylphenyl, 4-trifluoromethylphenyl, 4-(methoxymethyl)phenyl, 4-  
20 (2-hydroxyethyl)phenyl, 3-carboxylphenyl, 4-carboxylphenyl, 4-methoxyphenyl, 4-carbamoylphenyl, 4-methylthiophenyl, 4-(dimethylaminocarbonyl)phenyl, 4-methylsulfonylphenyl, benzyl, phenethyl, benzyloxy, 4-fluorobenzyloxy, 4-chlorobenzyloxy, 2-thiazolyl, 3-pyridyl, 4-pyridyl, 4-pyridylmethyloxy, 2-  
25 piperidylmethyloxy, 3-piperidylmethyloxy, 4-piperidylmethyloxy, 1-methylpiperidin-4-ylmethyloxy, 1-acetylpiperidin-4-ylmethyloxy, 2-chloropiperidin-4-ylmethyloxy, 1-(methylsulfonyl)piperidin-4-ylmethyloxy, 2-methylthiazol-4-yl, (2-methylthiazol-4-yl)methyloxy, (2,4-dimethylthiazol-5-yl)methyloxy, 5-tetrazolyl,  
30 3-fluorobenzoyl, piperidinocarbonyl, 4-hydroxypiperidinocarbonyl, 1-pyrrolidinylcarbonyl, morpholinocarbonyl, 4-thiomorpholinylcarbonyl, benzylaminocarbonyl, N-benzyl-N-methylaminocarbonyl, (4-pyridylmethyl)aminocarbonyl and (cyclohexylmethyl)aminocarbonyl.

35 Most preferable substituents are fluorine atom, chlorine atom, methyl, hydroxymethyl, carboxyl, carbamoyl, methylaminocarbonyl, dimethylaminocarbonyl, methoxy, methylamino, acetylamino, aminosulfonyl, dimethylaminosulfonyl, tert-

butylaminosulfonyl, phenyl, 3-fluorophenyl, 4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 3,5-dichlorophenyl, 4-methylphenyl, 4-tert-butylphenyl, 4-trifluoromethylphenyl, 4-carboxylphenyl, 4-methoxyphenyl, 4-carbamoylphenyl, 4-methylthiophenyl, 4-  
5 (dimethylaminocarbonyl)phenyl, 4-methylsulfonylphenyl and 2-oxopyrrolidin-1-yl.

The w is preferably 1 or 2, r and t are preferably 0, 1 or 2, particularly preferably 0 or 1, more preferably 0, p is preferably 1, and q is preferably 0 or 2.

10 In formula [I], when X is



wherein each symbol is as defined above and w is 2 or above, one of Z is preferably C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D or heterocyclic group  
15 optionally substituted by 1 to 5 substituent(s) selected from group D, particularly preferably C<sub>6-14</sub> aryl optionally substituted by 1 to 5 substituent(s) selected from group D.

The pharmaceutically acceptable salt may be any as long as it forms a non-toxic salt with a compound of the above-mentioned  
20 formula [I] or [II]. Such salt can be obtained by reacting the compound with an inorganic acid, such as hydrochloric acid, sulfuric acid, phosphoric acid, hydrobromic acid and the like, or an organic acid, such as oxalic acid, malonic acid, citric acid, fumaric acid, lactic acid, malic acid, succinic acid, tartaric  
25 acid, acetic acid, trifluoroacetic acid, gluconic acid, ascorbic acid, methylsulfonic acid, benzylsulfonic acid and the like, or an inorganic base, such as sodium hydroxide, potassium hydroxide, calcium hydroxide, magnesium hydroxide, ammonium hydroxide and the like, or an organic base, such as methylamine, diethylamine,  
30 triethylamine, triethanolamine, ethylenediamine, tris(hydroxymethyl)methylamine, guanidine, choline, cinchonine and the like, with an amino acid, such as lysine, arginine, alanine and the like. The present invention encompasses water-retaining product, hydrate and solvate of each compound.

35 The compounds of the above-mentioned formula [I] or [II] have various isomers. For example, E compound and Z compound are present as geometric isomers, and when the compound has an

asymmetric carbon, an enantiomer and a diastereomer are present due to the asymmetric carbon. A tautomer may be also present. The present invention encompasses all of these isomers and mixtures thereof.

5       The present invention also encompasses prodrug and metabolite of each compound.

A prodrug means a derivative of the compound of the present invention, which is capable of chemical or metabolic decomposition, which shows inherent efficacy by reverting to the  
10 original compound after administration to a body, and which includes salts and complexes without a covalent bond.

When the inventive compound is used as a pharmaceutical preparation, the inventive compound is generally admixed with pharmaceutically acceptable carriers, excipients, diluents,  
15 binders, disintegrators, stabilizers, preservatives, buffers, emulsifiers, aromatics, coloring agents, sweeteners, thickeners, correctives, solubilizers, and other additives such as water, vegetable oil, alcohol such as ethanol, benzyl alcohol and the like, polyethylene glycol, glycerol triacetate, gelatin, lactose,  
20 carbohydrate such as starch and the like, magnesium stearate, talc, lanolin, petrolatum and the like, and prepared into a dosage form of tablets, pills, powders, granules, suppositories, injections, eye drops, liquids, capsules, troches, aerosols, elixirs, suspensions, emulsions, syrups and the like, which can  
25 be administered systemically or topically and orally or parenterally.

While the dose varies depending on the age, body weight, general condition, treatment effect, administration route and the like, it is from 0.1 mg to 1 g for an adult per dose, which is  
30 given one to several times a day.

The prophylaxis of hepatitis C means, for example, administration of a pharmaceutical agent to an individual found to carry an HCV by a test and the like but without a symptom of hepatitis C, or to an individual who shows an improved disease  
35 state of hepatitis after a treatment of hepatitis C, but who still carries an HCV and is associated with a risk of recurrence of hepatitis.

Inasmuch as HCV is known to be a virus associated with many genetic mutations, a compound effective for many genotypes is one of the preferable modes. If a compound ensures high blood concentration when administered as a pharmaceutical agent to an animal infected with HCV, it is also one of the preferable modes. From these aspects, a compound having high inhibitory activity on both HCV type 1a and type 1b and high blood concentration, such as 2-{4-[2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride, is particularly preferable.

The fused ring compound of the formula [I] or [II] of the present invention can be administered to mammals inclusive of human for the purpose of prevention or treatment of hepatitis C or inhibition of hepatitis C virus polymerase. The fused ring compound of the present invention can be also administered to mammals inclusive of human along with at least one pharmaceutical agent (hereinafter combination drug) selected from an antiviral agent other than the compound of the formula [I], an antiinflammatory agent and an immunostimulant for the purpose of prevention or treatment of hepatitis C or inhibition of hepatitis C virus polymerase. In the case of combined administration, the compound of the present invention can be administered simultaneously with the combination drug or administered at certain time intervals. In the case of combined administration, a pharmaceutical composition containing the compound of the present invention and a combination drug can be administered. Alternatively, a pharmaceutical composition containing the compound of the present invention and a pharmaceutical composition containing a combination drug may be administered separately. The administration route may be the same or different.

In the case of a combined administration, the compound of the present invention can be administered once a day or several times a day in a single dose of 0.1 mg to 1 g, or may be administered in a smaller dose. The combination drug can be administered in a dose generally used for the prevention or treatment of hepatitis C or in a smaller dose.



Examples of other antiviral agent include interferons (interferon  $\alpha$ , interferon  $\beta$ , interferon  $\gamma$  etc.), Ribavirin (1- $\beta$ -D-ribofuranosyl-1H-1,2,4-triazole-3-carboxamide) and the like.

Examples of the production method of the compound to be used for the practice of the present invention are given in the following. However, the production method of the compound of the present invention is not limited to these examples.

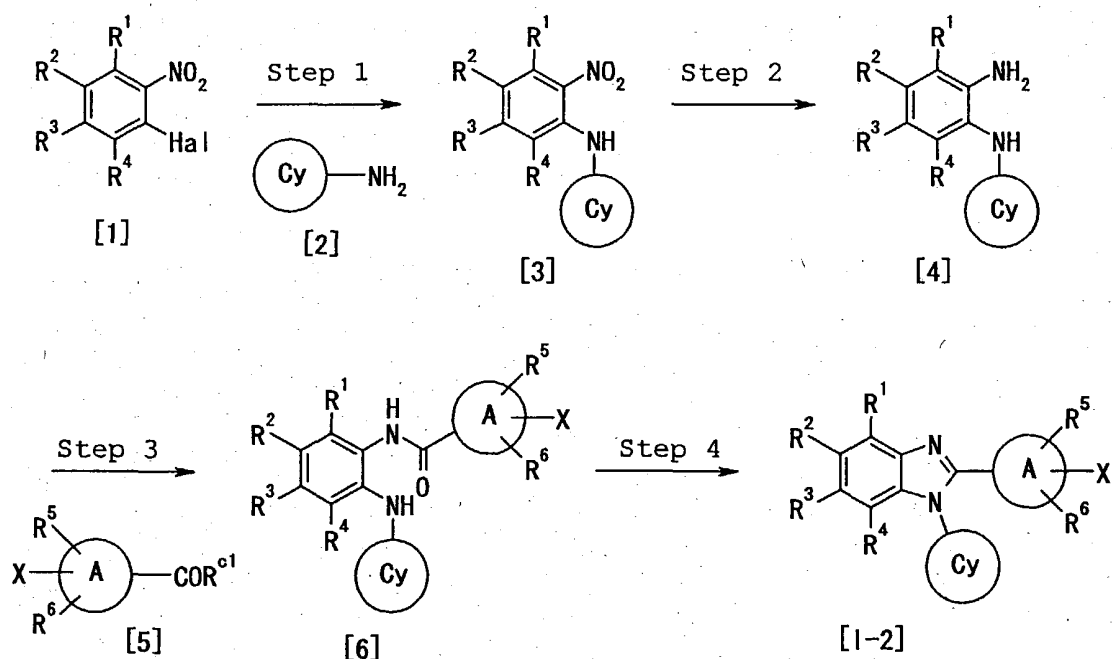
Even if no directly corresponding disclosure is found in the following Production Methods, the steps may be modified for efficient production of the compound, such as introduction of a protecting group into a functional group with deprotection in a subsequent step, and changing the order of Production Methods and steps.

The treatment after reaction in each step may be conventional ones, for which typical methods, such as isolation and purification, crystallization, recrystallization, silica gel chromatography, preparative HPLC and the like, can be appropriately selected and combined.

#### Production Method 1

In this Production Method, a benzimidazole compound is formed from a nitrobenzene compound.

#### Production Method 1-1



wherein  $Hal$  is halogen atom, such as chlorine atom, bromine atom and the like,  $R^{c1}$  is halogen atom, such as chlorine atom, bromine

atom and the like, or hydroxyl group, and other symbols are as defined above.

#### Step 1

A compound [1] obtained by a conventional method or a commercially available compound [1] is reacted with amine compound [2] in a solvent such as N,N-dimethylformamide (DMF), acetonitrile, tetrahydrofuran (THF), toluene and the like in the presence or absence of a base such as potassium carbonate, triethylamine, potassium t-butoxide and the like at room temperature or with heating to give compound [3].

#### Step 2

The compound [3] is hydrogenated in a solvent such as methanol, ethanol, THF, ethyl acetate, acetic acid, water and the like in the presence of a catalyst such as palladium carbon, palladium hydroxide, platinum oxide, Raney nickel and the like at room temperature or with heating to give compound [4]. In addition, compound [3] is reduced with a reducing agent such as zinc, iron, tin(II) chloride, sodium sulfite and the like, or reacted with hydrazine in the presence of iron(III) chloride to give compound [4].

#### Step 3

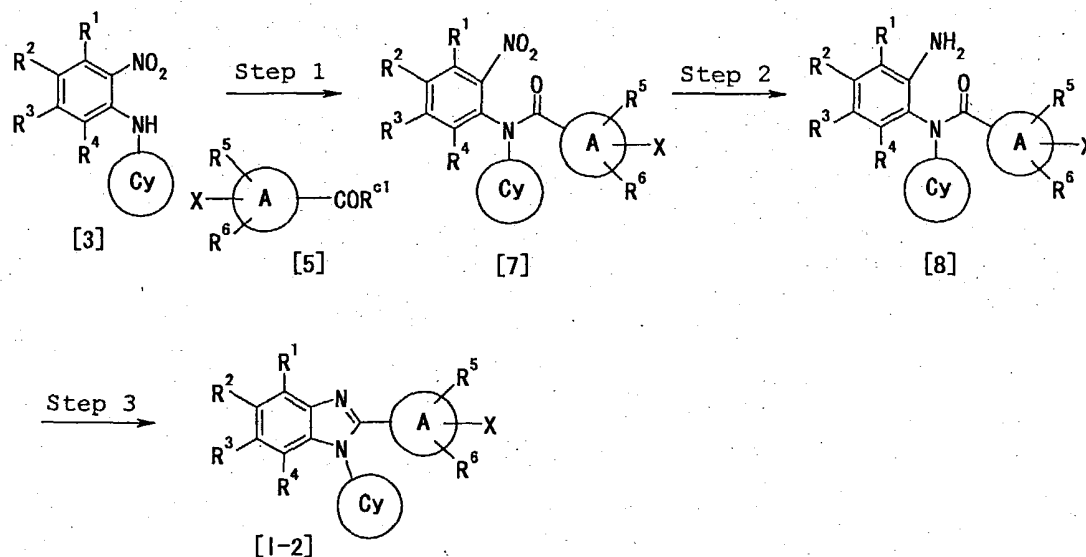
The compound [4] is condensed with carboxylic acid compound [5] in a solvent such as DMF, acetonitrile, THF, chloroform, ethyl acetate, methylene chloride, toluene and the like using a condensing agent such as dicyclohexylcarbodiimide, 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride, diphenylphosphoryl azide and the like and, where necessary, adding N-hydroxysuccinimide, 1-hydroxybenzotriazole and the like to give amide compound [6]. Alternatively, amide compound [6] can be obtained from compound [5] as follows. The carboxylic acid compound [5] is converted to an acid halide derived with thionyl chloride, oxalyl chloride and the like, or an active ester (e.g., mixed acid anhydride derived with ethyl chlorocarbonate and the like), which is then reacted in the presence of a base, such as triethylamine, potassium carbonate, pyridine and the like, or in an amine solvent, such as pyridine and the like, to give amide compound [6].

#### Step 4

The compound [6] is heated in a solvent such as ethanol, methanol, toluene, DMF, chloroform and the like or without a solvent in the presence of an acid such as acetic acid, formic acid, hydrochloric acid, dilute sulfuric acid, phosphoric acid, polyphosphoric acid, p-toluenesulfonic acid and the like, a halogenating agent such as zinc chloride, phosphorus oxychloride, thionyl chloride and the like or acid anhydride such as acetic anhydride and the like, to allow cyclization to give compound [I-2].

#### 10 Production Method 1-2

This Production Method is an alternative method for producing compound [I-2].



wherein each symbol is as defined above.

#### 15 Step 1

The compound [3] obtained in the same manner as in Step 1 of Production Method 1-1 is subjected to amide condensation with compound [5] in the same manner as in Step 3 of Production Method 1-1 to give compound [7].

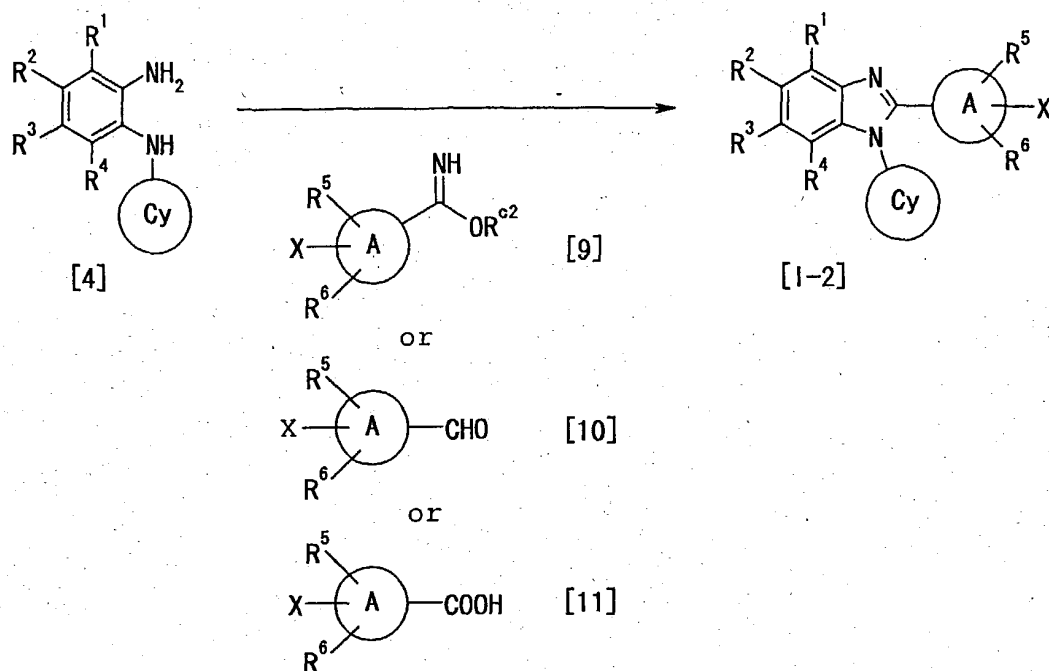
#### 20 Step 2

The compound [7] is reduced in the same manner as in Step 2 of Production Method 1-1 to give compound [8].

#### Step 3

The compound [8] is subjected to cyclization in the same manner as in Step 4 of Production Method 1-1 to give compound [I-2].

#### Production Method 1-3



wherein  $R^{c2}$  is alkyl such as methyl, ethyl and the like, and other symbols are as defined above.

The compound [4] is reacted with imidate compound [9] in a solvent such as methanol, ethanol, acetic acid, DMF, THF, chloroform and the like at room temperature or with heating to give compound [I-2].

In addition, compound [4] may be reacted with aldehyde compound [10] in a solvent such as acetic acid, formic acid, acetonitrile, DMF, nitrobenzene, toluene and the like in the presence or absence of an oxidizing agent such as benzofuroxan, manganese dioxide, 2,3-dichloro-5,6-dicyano-p-benzoquinone, iodine, potassium ferricyanide and the like with heating to give compound [I-2].

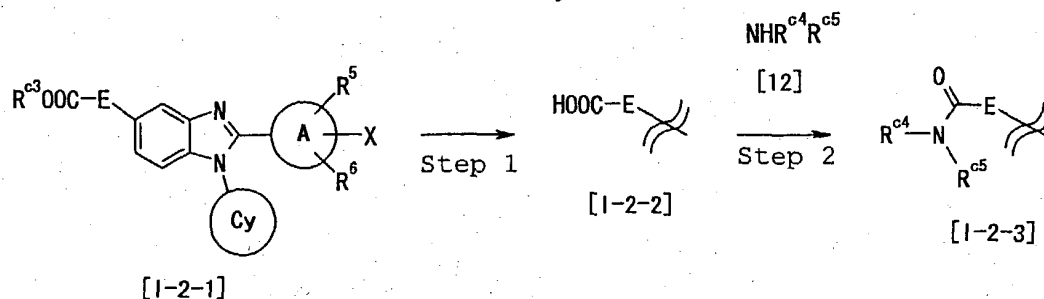
Alternatively, compound [4] and carboxylic acid compound [11] may be heated to allow reaction in the presence of polyphosphoric acid, phosphoric acid, phosphorus oxychloride, hydrochloric acid and the like to give compound [I-2].

#### Production Method 2

In this Production Method, conversion of the substituents ( $R^1, R^2, R^3, R^4$ ) on the benzene ring of benzimidazole is shown. While a method of converting  $R^2$  when  $R^1, R^3$  and  $R^4$  are hydrogen atoms is shown, this Production Method is applicable irrespective of the position of substitution.

## Production Method 2-1

Conversion of carboxylic acid ester moiety to amide



wherein E is a single bond,  $-(\text{CH}_2)_s-$ ,  $-\text{O}-(\text{CH}_2)_s-$  or  $-\text{NH}-(\text{CH}_2)_s-$  (wherein s is an integer of 1 to 6),  $\text{R}^{\text{c}3}$ ,  $\text{R}^{\text{c}4}$  and  $\text{R}^{\text{c}5}$  are  $\text{C}_{1-6}$  alkyl, and other symbols are as defined above.

### Step 1

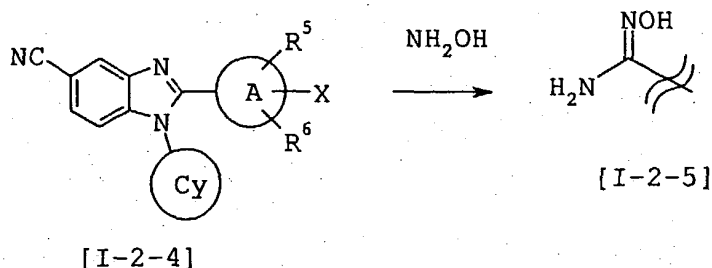
The compound [I-2-1] obtained in the same manner as in the above-mentioned Production Method is subjected to hydrolysis in a solvent such as methanol, ethanol, THF, dioxane and the like, or in a mixed solvent of these solvents and water under basic conditions with sodium hydroxide, potassium hydroxide, potassium carbonate, lithium hydroxide and the like or under acidic conditions with hydrochloric acid, sulfuric acid and the like to give compound [I-2-2].

### Step 2

The compound [I-2-2] is reacted with compound [12] in the same manner as in Step 3 of Production Method 1-1 to give compound [I-2-3].

## 20 Production Method 2-2

Conversion of cyano group to substituted amidino group



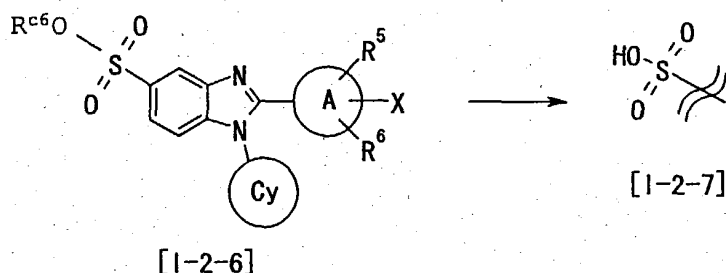
wherein each symbol is as defined above.

The compound [I-2-4] obtained in the same manner as in the above-mentioned Production Method is reacted with hydroxylamine in a solvent such as water, methanol, ethanol, THF, DMF and the like to give compound [I-2-5]. When a salt of hydroxylamine such

as hydrochloride and the like is used, the reaction is carried out in the presence of a base such as sodium hydrogencarbonate, sodium hydroxide, triethylamine and the like.

### Production Method 2-3

- 5 Conversion of sulfonic acid ester moiety to sulfonic acid



wherein  $R^6$  is  $C_{1-6}$  alkyl, and other symbols are as defined above.

- The compound [I-2-6] obtained in the same manner as in the above-mentioned Production Method is reacted with iodide salt such as sodium iodide, lithium iodide and the like, bromide salt such as sodium bromide, trimethylammonium bromide and the like, amine such as pyridine, trimethylamine, triazole and the like, phosphine such as triphenylphosphine and the like in a solvent such as DMF, dimethyl sulfoxide (DMSO), acetonitrile, methanol, ethanol, water and the like with heating to give compound [I-2-7].

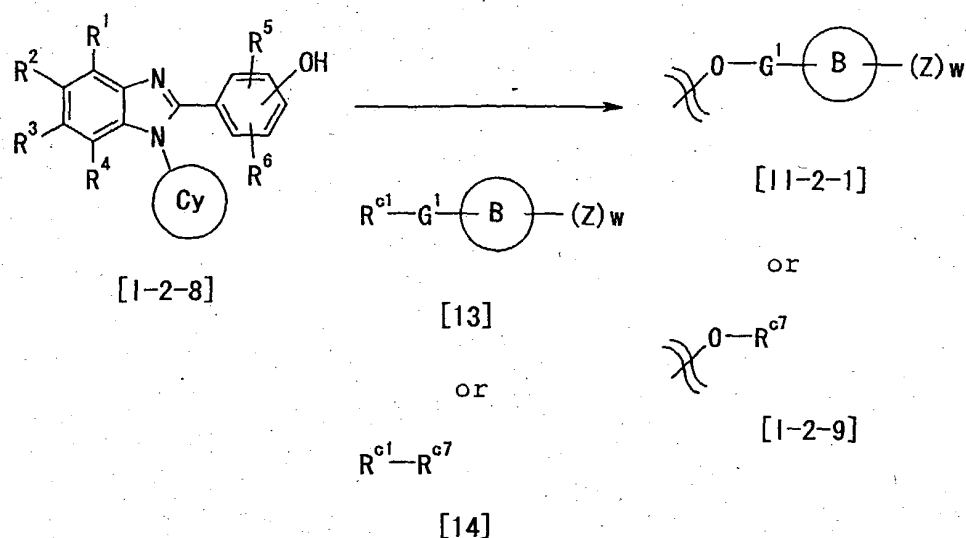
### Production Method 3

This Production Method relates to conversion of the substituent(s) on phenyl group at the 2-position of benzimidazole.

- 20 This Production Method can be used even when phenyl is a different ring.

### Production Method 3-1

Conversion of hydroxyl group to ether



wherein  $R^{c7}$  is optionally substituted alkyl corresponding to  $R^{a11}$ ,  $G^1$  is a single bond,  $-(CH_2)_n-$ ,  $-(CH_2)_n-O-$ ,  $-(CH_2)_n-CO-$  or  $-(CH_2)_m-CR^{a15}R^{a16}-(CH_2)_n-$ , wherein \* show the side to be bonded to  $R^{c1}$ , and other symbols are as defined above.

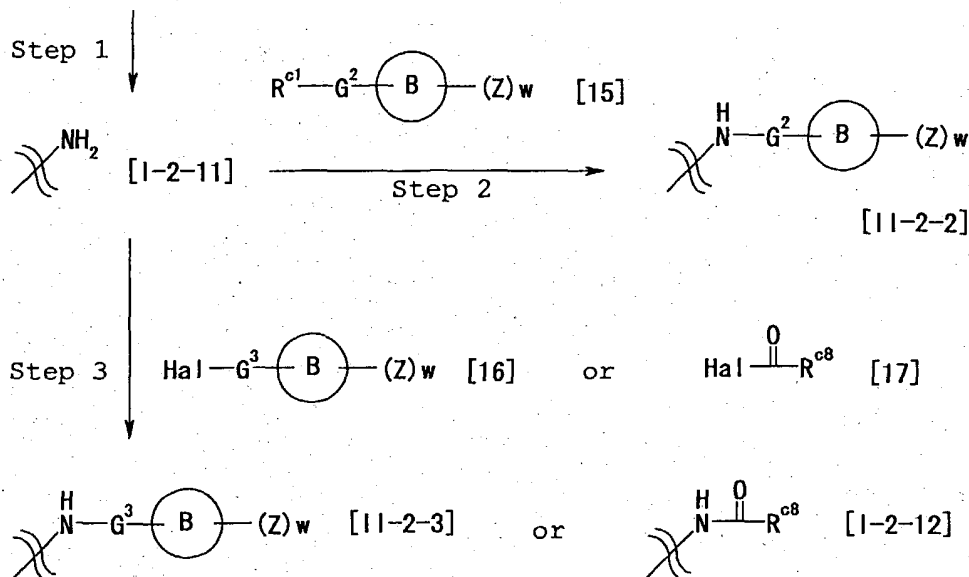
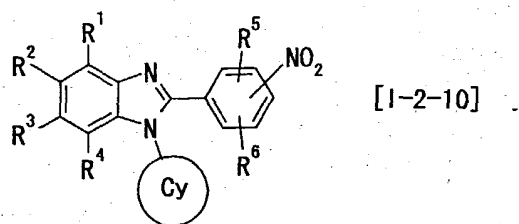
When  $R^{c1}$  of compound [13] is halogen atom, compound [I-2-8] obtained in the same manner as in the above-mentioned Production Method is reacted with compound [13] in a solvent such as DMF, DMSO, acetonitrile, ethanol, THF and the like in the presence of a base such as sodium hydride, sodium hydroxide, potassium hydroxide, potassium carbonate, sodium ethoxide, potassium t-butoxide and the like at room temperature or with heating to give compound [II-2-1].

When  $R^{c1}$  of compound [13] is hydroxyl group, the hydroxyl group of compound [13] is converted to halogen atom with thionyl chloride, phosphorus tribromide, carbon tetrabromide - triphenylphosphine and the like and reacted with compound [I-2-8] by the aforementioned method to give compound [II-2-1]. In this case, compound [I-2-8] may be subjected to Mitsunobu reaction with compound [13] in a solvent such as DMF, acetonitrile, THF and the like using triphenylphosphine - diethyl azodicarboxylate and the like to give compound [II-2-1].

The compound [I-2-9] can be obtained in the same manner from compound [I-2-8] and compound [14].

## 25 Production Method 3-2

Conversion of nitro to substituted amino group



wherein  $R^{c8}$  is  $C_{1-6}$  alkyl,  $G^2$  is  $^*-(CH_2)_n-$  or  $^*-\text{CHR}^{a15}$ ,  $G^3$  is  $-\text{CO}-$ ,  $^*-\text{CO}_2-$ ,  $^*-\text{CONH}-$  or  $-\text{SO}_2-$ , and other symbols are as defined above.

#### Step 1

5 The nitro compound [I-2-10] obtained in the same manner as in the above-mentioned Production Method is reacted in the same manner as in Step 2 of Production Method 1-1 to give compound [I-2-11].

#### Step 2

10 The compound [I-2-11] is alkylated with compound [15] in the same manner as in Production Method 3-1 to give compound [II-2-2].

#### Step 3

When  $G^3$  of compound [16] is  $-\text{CO}-$ ,  $-\text{CO}_2-$  or  $-\text{CONH}-$ , compound [I-2-11] is acylated with compound [16] in the same manner as in  
15 Step 3 of Production Method 1-1 to give compound [II-2-3].

When  $G^3$  of compound [16] is  $-\text{SO}_2-$ , sulfonylation is conducted using sulfonyl halide instead of acid halide used in Step 3 of Production Method 1-1 to give compound [II-2-3].

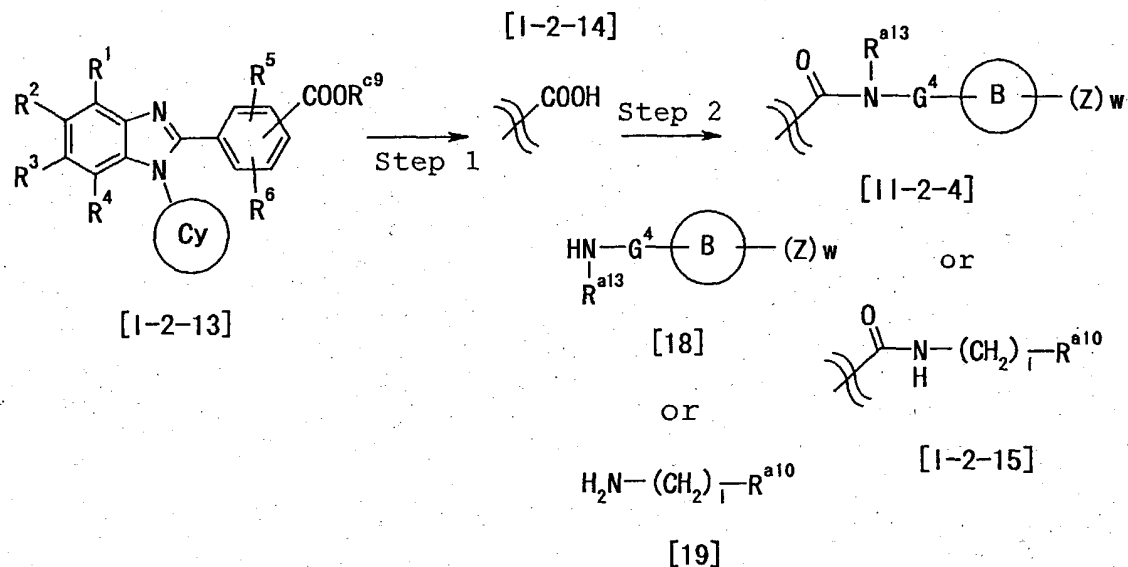
The compound [I-2-11] is acylated with compound [17] in the  
20 same manner as above to give compound [I-2-12].



This Production Method is applied in the same manner as above to give disubstituted compounds (tertiary amine) of compound [II-2-2], compound [II-2-3] and compound [I-2-12].

### Production Method 3-3

#### 5 Conversion of carboxylic acid ester moiety to amide



wherein R<sup>c9</sup> is C<sub>1-6</sub> alkyl, G<sup>4</sup> is #-(CH<sub>2</sub>)<sub>n</sub>-, #-(CH<sub>2</sub>)<sub>n</sub>-NH- or #-CHR<sup>a14</sup>- wherein # shows the side that is bounded to amine and other

10 symbols are as defined above.

#### Step 1

The compound [I-2-13] obtained in the same manner as in the above-mentioned Production Method is reacted in the same manner as in Step 1 of Production Method 2-1 to give compound [I-2-14].

#### Step 2

The compound [I-2-14] is reacted with compound [18] in the same manner as in Step 2 of Production Method 2-1 to give compound [II-2-4].

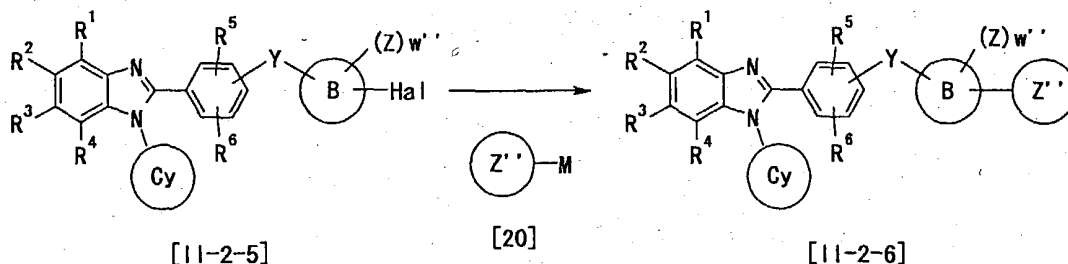
20 The compound [I-2-15] is obtained from compound [I-2-14] and compound [19] in the same manner as above.

### Production Method 4

In this Production Method, additional substituent(s) is(are) introduced into ring B on phenyl group that substitutes the 2-position of benzimidazole. This Production Method is applicable even when phenyl is a different ring.

#### Production Method 4-1

Direct bonding of ring Z'' to ring B



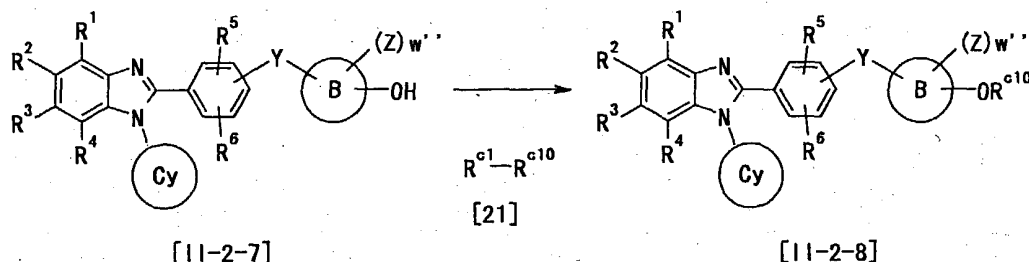
wherein ring Z''-M is aryl metal compound, ring Z'' moiety is optionally substituted C<sub>6-14</sub> aryl or optionally substituted

5 heterocyclic group corresponding to substituent Z, and the metal moiety contains boron, zinc, tin, magnesium and the like, such as phenylboronic acid, w'' is 0, 1 or 2, and other symbols are as defined above.

The compound [II-2-5] obtained in the same manner as in the  
 10 above-mentioned Production Method is reacted with aryl metal compound [20] in a solvent such as DMF, acetonitrile, 1,2-dimethoxyethane, THF, toluene, water and the like in the presence of a palladium catalyst such as tetrakis(triphenylphosphine)-palladium, bis(triphenylphosphine)palladium(II) dichloride,  
 15 palladium acetate - triphenylphosphine and the like, a nickel catalyst such as nickel chloride, [1,3-bis(diphenylphosphino)-propane]nickel(II) chloride and the like, and a base such as potassium carbonate, potassium hydrogencarbonate, sodium hydrogen-carbonate, potassium phosphate, triethylamine and the like at room  
 20 temperature or with heating, to give compound [II-2-6].

#### Production Method 4-2

Conversion of hydroxyl group to ether



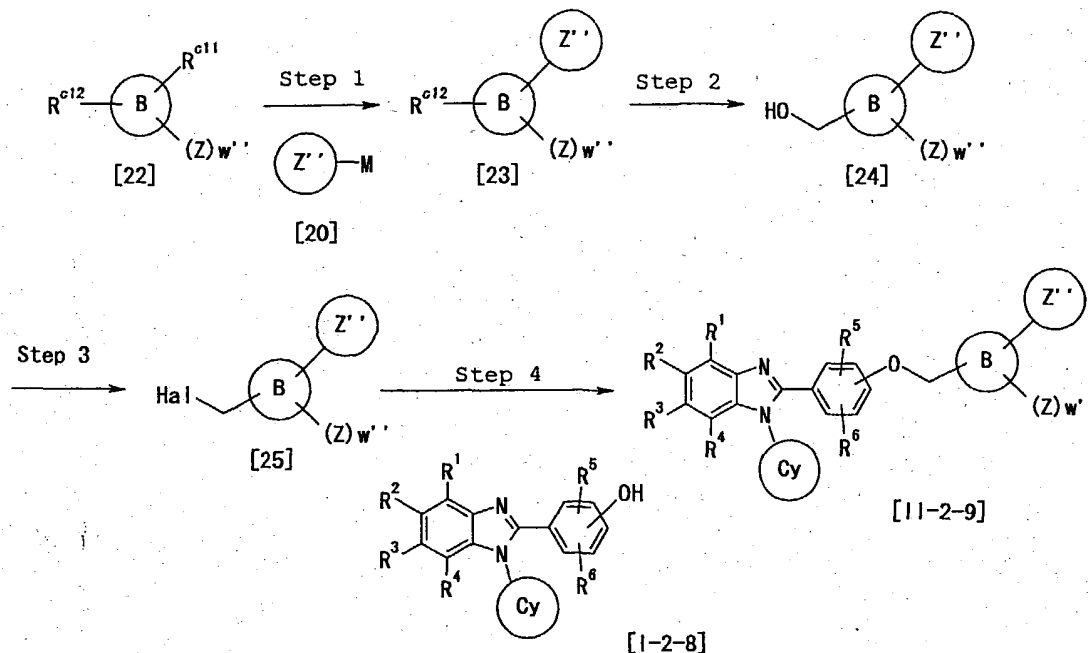
wherein R<sup>c10</sup> is -R<sup>a20</sup> or -(CH<sub>2</sub>)<sub>p</sub>-COR<sup>a21</sup> corresponding to substituent  
 25 Z, and other symbols are as defined above.

The compound [II-2-7] obtained in the same manner as in the above-mentioned Production Method is reacted with compound [21]

in the same manner as in Production Method 3-1 to give compound [II-2-8].

### Production Method 4-3

Synthesis in advance of ring B part such as compound [13] in  
5 Production Method 3-1



wherein  $R^{c11}$  is leaving group such as bromine atom, iodine atom, trifluoromethanesulfonyloxy and the like,  $R^{c12}$  is formyl, carboxyl  
10 or carboxylic acid ester such as methoxycarbonyl, ethoxycarbonyl, tert-butoxycarbonyl and the like, and other symbols are as defined above.

#### Step 1

Commercially available compound [22] or compound [22]  
15 obtained by a conventional method is reacted with aryl metal compound [20] in the same manner as in Production Method 4-1 to give compound [23].

#### Step 2

The compound [23] obtained in the same manner as in the  
20 above-mentioned Production Method is reduced according to a conventional method to give compound [24].

For example, compound [23] is reacted with in a solvent such as methanol, ethanol, THF and the like in the presence of a reducing agent such as lithium aluminum hydride, sodium

borohydride and the like under cooling to heating to give compound [24].

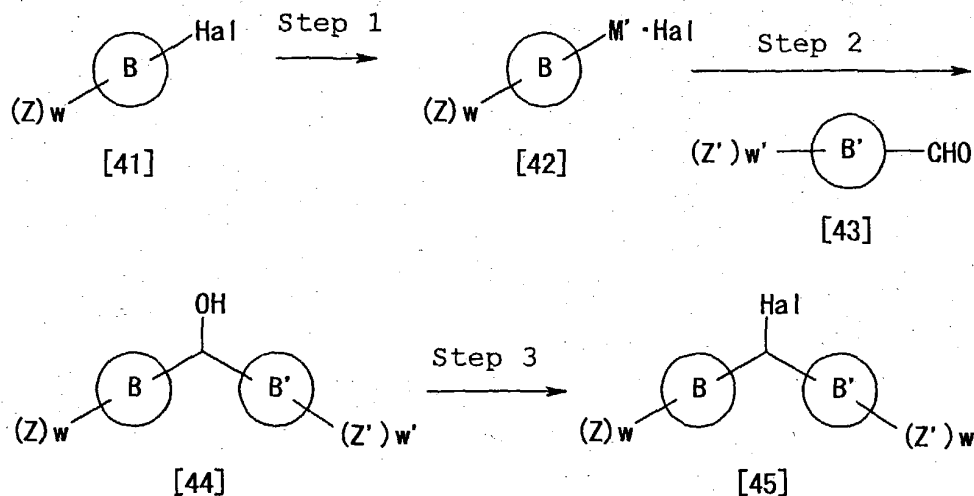
### Step 3

The compound [24] obtained in the same manner as in the above-mentioned Production Method is reacted in a solvent such as 1,4-dioxane, diethyl ether, THF, dichloromethane, chloroform, toluene and the like with a halogenating agent, such as phosphorus pentachloride, phosphorus tribromide, thionyl chloride and the like, in the presence of a tertiary amine such as pyridine and the like to give compound [25].

### Step 4

The compound [24] or [25] obtained in the same manner as in the above-mentioned Production Method is reacted with compound [I-2-8] in the same manner as in Production Method 3-1 to give compound [II-2-9].

### Production Method 4-4



wherein M' is a metal such as magnesium, lithium, zinc and the like, and other symbols are as defined above.

### Step 1

Commercially available compound [41] or compound [41] obtained by a conventional method is converted to aryl metal reagent by a conventional method to give compound [42].

For example, when M' is magnesium, magnesium is reacted with compound [41] in a solvent such as THF, diethyl ether, benzene, toluene and the like, preferably THF, from cooling to heating preferably at -100°C to 100°C to give compound [42].

### Step 2

The compound [42] obtained in the same manner as in the above-mentioned Production Method is reacted with compound [43] to give compound [44].

The compound [42] is reacted in a solvent such as diethyl  
5 ether, benzene, toluene, THF and the like, preferably THF, from cooling to room temperature, preferably at -100°C to 30°C to give compound [44].

### Step 3

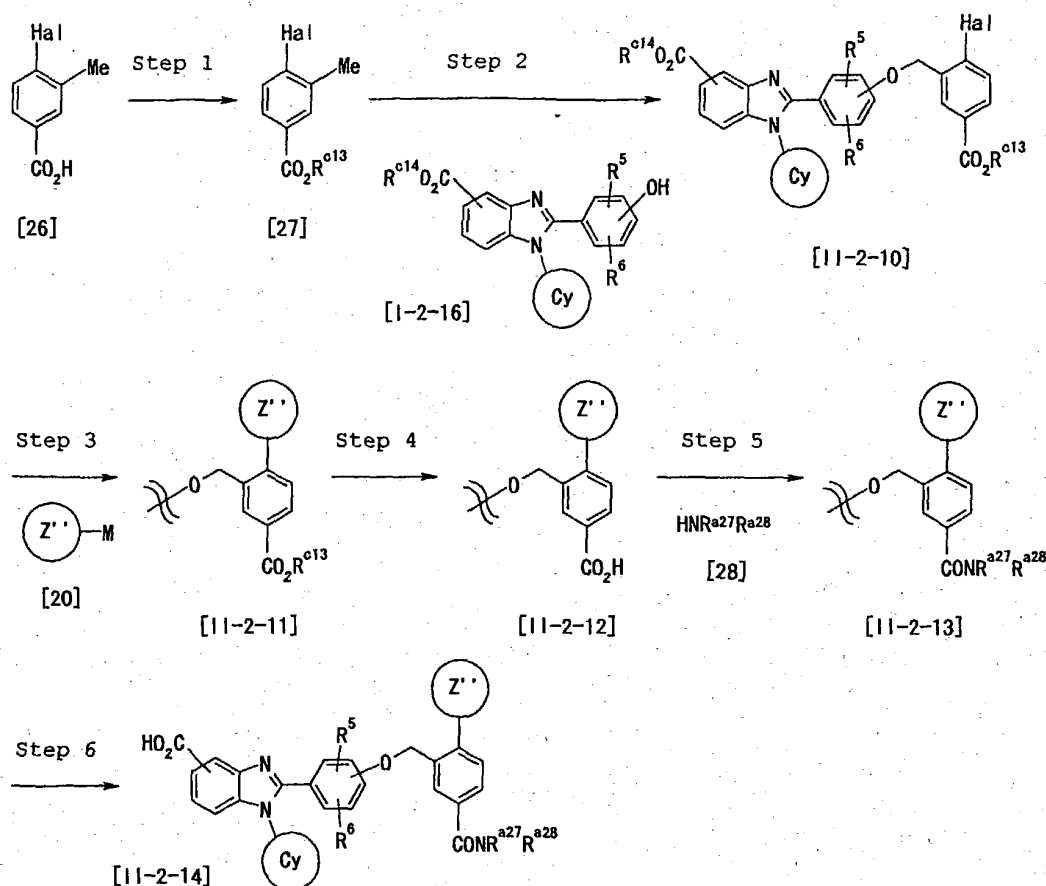
The compound [44] obtained in the same manner as in the  
10 above-mentioned Production Method is halogenated in the same manner as in Step 3 of Production Method 4-3 to give compound [45].

The compound [44] is reacted with thionyl chloride and pyridine preferably in toluene solvent to give compound [45].

15 When compound [45] is symmetric, namely, when the ring B-(Z)w moiety and the ring B'-(Z')w' moiety are the same, compound [42] is reacted with formate such as methyl formate, ethyl formate and the like, preferably ethyl formate, in a solvent such as diethyl ether, benzene, toluene, THF and the like, preferably  
20 THF, from cooling to room temperature, preferably at -100°C to 30°C, to give compound [45].

### Production Method 4-5

Method including steps to introduce a protecting group into a functional group



wherein  $\text{R}^{\text{c}13}$  is carboxylic acid protecting group such as tert-butyl and the like,  $\text{R}^{\text{c}14}$  is carboxylic acid protecting group such as methyl and the like and other symbols are as defined above.

### Step 1

Commercially available compound [26] or compound [26] obtained by a conventional method is protected by a conventional method to give compound [27].

For example, when  $\text{R}^{\text{c}13}$  is tert-butyl, compound [26] is converted to acid halide with thionyl chloride, oxalyl chloride and the like in a solvent such as THF, chloroform, dichloromethane, toluene and the like, and reacted with potassium tert-butoxide to give compound [27].

As used herein,  $\text{R}^{\text{c}13}$  may be a different protecting group as long as it is not removed during the Step 2 or Step 3 but removed in Step 4 without affecting  $-\text{CO}_2\text{R}^{\text{c}14}$ .

### Step 2

The methyl group of compound [27] obtained in the same manner as in the above-mentioned Production Method is converted

to bromomethyl with N-bromosuccinimide and N,N'-azobisisobutyronitrile and reacted with compound [I-2-16] in the same manner as in Production Method 3-1 to give compound [II-2-10].

5 **Step 3**

The compound [II-2-10] obtained in the same manner as in the above-mentioned Production Method is reacted with aryl metal compound [20] in the same manner as in Production Method 4-1 to give compound [II-2-11].

10 **Step 4**

The  $R^{c13}$  of the compound [II-2-11] obtained in the same manner as in the above-mentioned Production Method is removed by a conventional method to give compound [II-2-12].

The protecting group of carboxylic acid can be removed by a  
15 conventional deprotection method according to the protecting group. In this Step, the conditions free from reaction of  $R^{c14}$  are preferable. For example, when  $R^{c13}$  is tert-butyl, compound [II-2-11] is treated with trifluoroacetic acid in a solvent such as dichloromethane, chloroform and the like to give compound [II-2-  
20 12].

**Step 5**

The compound [II-2-12] obtained in the same manner as in the above-mentioned Production Method is subjected to amide condensation with compound [28] in the same manner as in Step 3  
25 of Production Method 1-1 to give compound [II-2-13].

**Step 6**

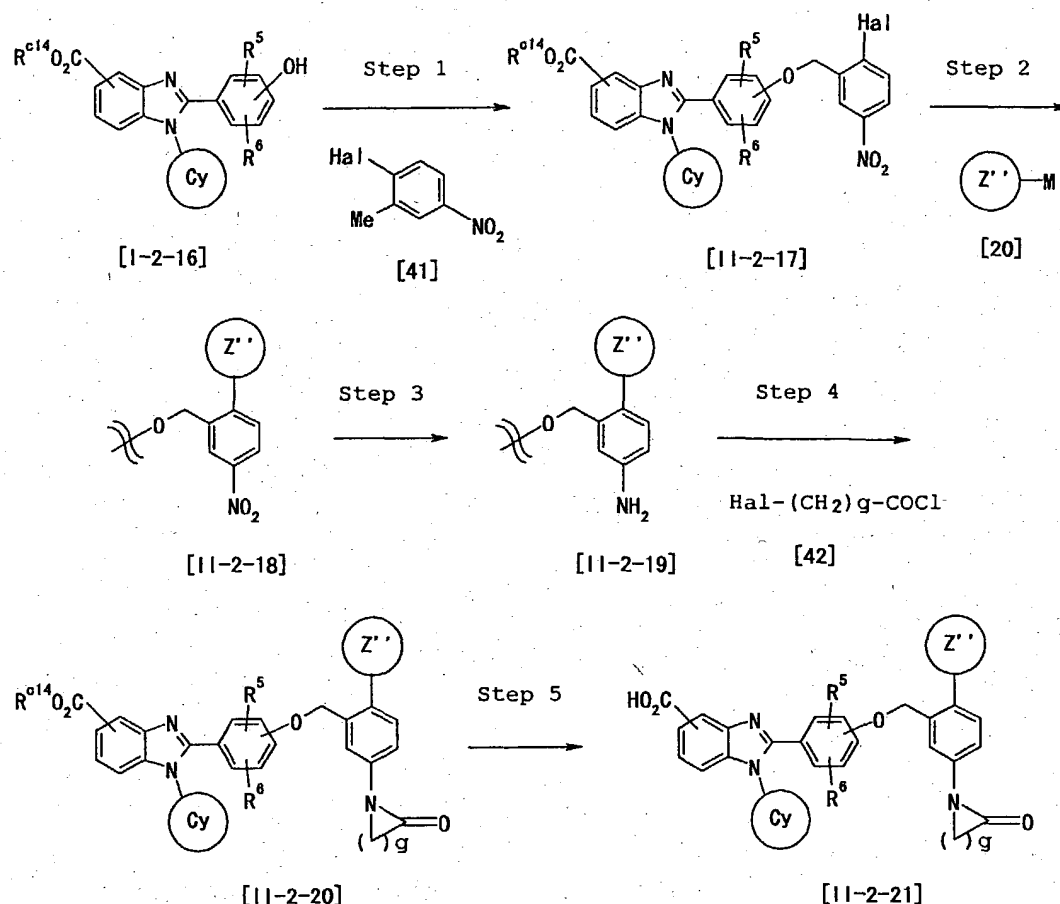
The compound [II-2-13] obtained in the same manner as in the above-mentioned Production Method is deprotected in the same manner as in Step 1 of Production Method 2-1 to give compound  
30 [II-2-14].

As used herein,  $R^{c14}$  is preferably a protecting group that does not react during the Step 1 through Step 5 but removed in this Step.

For example, when  $R^{c14}$  is methyl, compound [II-2-13] is  
35 reacted in an alcohol solvent such as methanol, ethanol, n-propanol, isopropanol and the like or a mixed solvent of alcohol solvent and water in the presence of a base such as potassium carbonate, sodium carbonate, lithium hydroxide, sodium hydroxide,

potassium hydroxide and the like from cooling to heating for deprotection, followed by acidifying the reaction solution to give compound [II-2-14].

#### Production Method 4-6



5

wherein  $g$  is an integer of 1 to 5, and other symbols are as defined above.

#### Step 1

The compound [I-2-16] obtained by the above-mentioned  
 10 Production Method is reacted with toluene derivative [41] in the same manner as in Step 2 of Production Method 4-5 to give compound [II-2-17].

#### Step 2

The compound [II-2-17] obtained by the above-mentioned  
 15 Production Method is reacted with aryl metal compound [20] in the same manner as in Production Method 4-1 to give compound [II-2-18].

#### Step 3



The compound [II-2-18] obtained by the above-mentioned Production Method is reduced in the same manner as in Step 2 of Production Method 1-1 to give compound [II-2-19].

#### Step 4

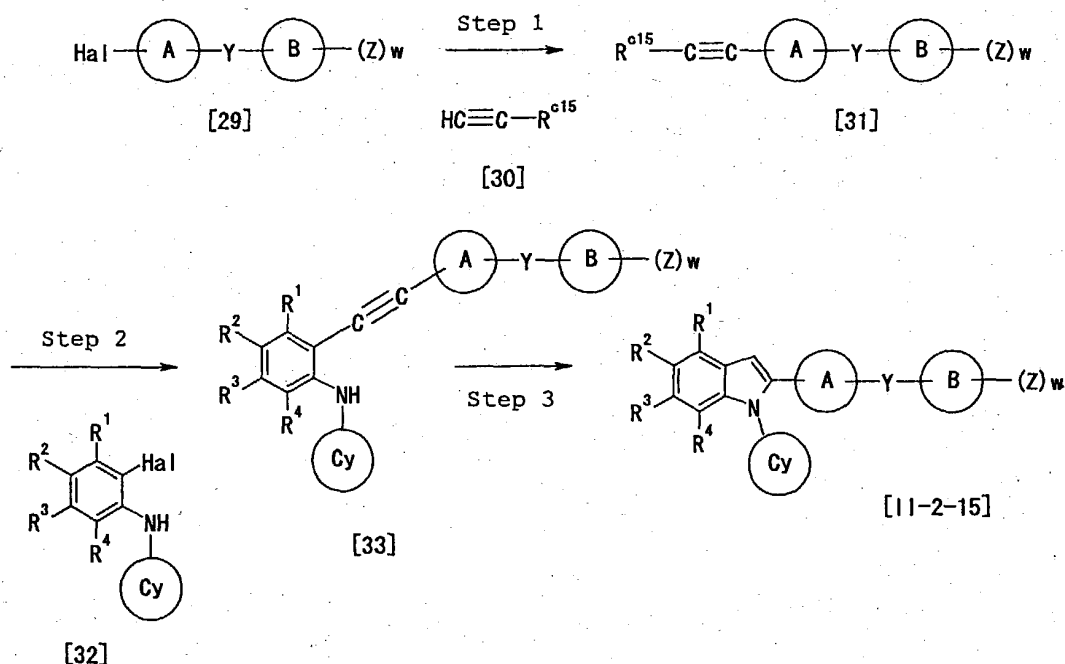
5 The compound [II-2-19] obtained by the above-mentioned Production Method is amide condensed with compound [42] in the same manner as in Step 3 of Production Method 1-1 and subjected to cyclization in the same manner as in Step 1 of Production Method 1-1 to give compound [II-2-20].

#### 10 Step 5

The compound [II-2-20] obtained by the above-mentioned Production Method is hydrolyzed in the same manner as in Step 1 of Production Method 2-1 to give compound [II-2-21].

#### Production Method 5

##### 15 Formation of indole ring



wherein  $\text{R}^{\text{C}15}$  is protecting group such as trimethylsilyl, tert-butyldimethylsilyl, tert-butyldiphenylsilyl and the like, and other symbols are as defined above.

#### 20 Step 1

The compound [29] obtained in the same manner as in the above-mentioned Production Method or conventional method is reacted with compound [30] in a solvent such as DMF, acetonitrile, 1,2-dimethoxyethane, THF, toluene, water and the like using a  
25 palladium catalyst such as tetrakis(triphenylphosphine)palladium,

bis(triphenylphosphine)palladium(II) dichloride, palladium acetate - triphenylphosphine and the like, a copper catalyst such as copper(I) iodide and the like or a mixture thereof, and in the presence of a base such as potassium carbonate, potassium hydrogencarbonate, sodium hydrogencarbonate, potassium phosphate, triethylamine and the like to give compound [31].

### Step 2

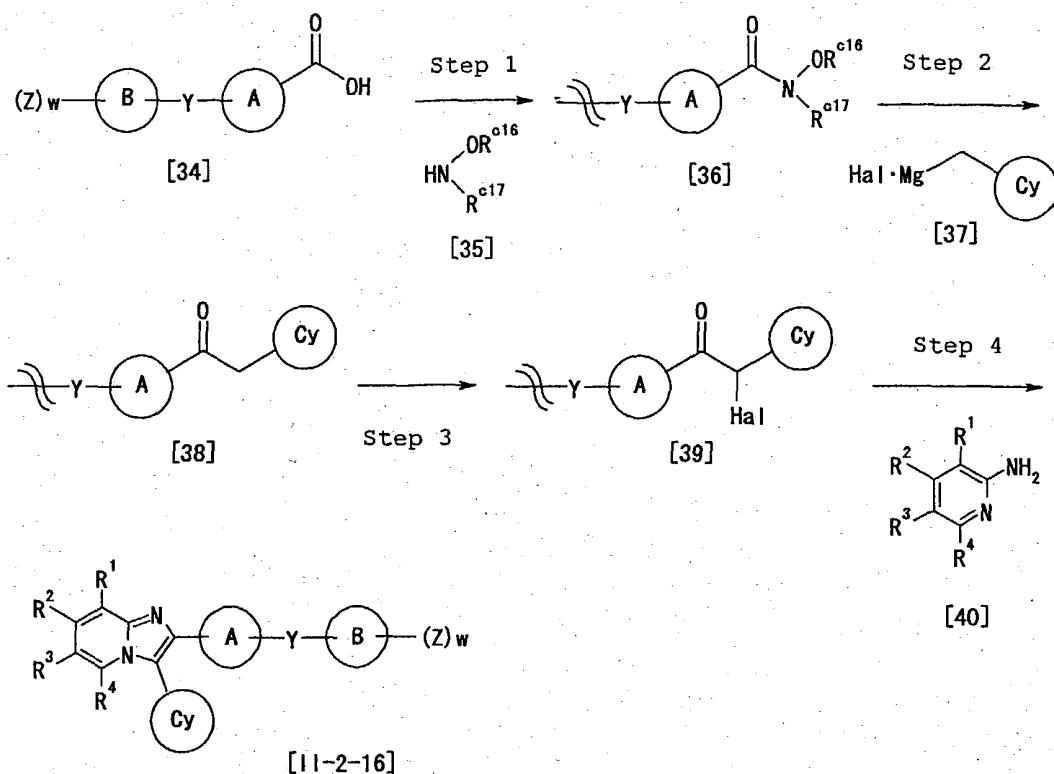
The compound [31] obtained in the same manner as in the above-mentioned Production Method is reacted in an alcohol solvent such as methanol, ethanol and the like or a mixed solvent of an alcohol solvent and a solvent such as DMF, acetonitrile, THF, chloroform, dichloromethane, ethyl acetate, methylene chloride, toluene and the like in the presence of a base such as potassium carbonate, sodium carbonate, lithium hydroxide, sodium hydroxide, potassium hydroxide, lithium hydride, sodium hydride, potassium hydride and the like at room temperature or with heating for deprotection, and reacted with compound [32] obtained in the same manner as in Step 1 of Production Method 1-1 in the same manner as in Step 1 of Production Method 5 to give compound [33].

### Step 3

The compound [33] obtained in the same manner as in the above-mentioned Production Method was subjected to cyclization in a solvent such as DMF, acetonitrile, THF, chloroform, dichloromethane, ethyl acetate, methylene chloride, toluene and the like in the presence of a copper catalyst such as copper(I) iodide and the like or a palladium catalyst such as palladium(II) chloride and the like at room temperature or with heating to give compound [II-2-15].

### 30 Production Method 6

Formation of imidazo[1,2-a]pyridine ring



wherein  $R^{c16}$  and  $R^{c17}$  are each independently alkyl, such as methyl, ethyl and the like, and other symbols are as defined above.

#### Step 1

- 5 The compound [34] obtained by the above-mentioned Production Method or a conventional method is subjected to amide condensation with compound [35] in the same manner as in Step 3 of Production Method 1-1 to give compound [36].

#### Step 2

- 10 The compound [36] obtained by the above-mentioned Production Method is reacted with Grignard reagent [37] obtained by a conventional method to give compound [38].

Alternatively, an acid halide of compound [34] may be used instead of compound [36].

#### 15 Step 3

The compound [38] obtained by the above-mentioned Production Method is subjected to halogenation by a conventional method to give compound [39].

- For example, when Hal is a bromine atom, compound [38] is  
 20 reacted with bromine under cooling or at room temperature in a solvent such as DMF, acetonitrile, THF, chloroform,

dichloromethane, ethyl acetate, toluene and the like to give compound [39].

Alternatively, a halogenating agent such as hypohalite (e.g., hypochlorite and the like), N-bromosuccinimide and the like may be used instead of bromine for halogenation.

#### Step 4

The compound [39] obtained by the above-mentioned Production Method is subjected to cyclization with compound [40] obtained by a conventional or known method (JP-A-8-48651) in the presence of a base such as potassium carbonate, sodium carbonate, lithium hydroxide, sodium hydroxide, potassium hydroxide, lithium hydride, sodium hydride, potassium hydride and the like in a solvent or without a solvent at room temperature or with heating to give compound [II-2-16].

In the compounds of the formulas [I] and [II], a desired heterocyclic group can be formed according to a method similar to the methods disclosed in known publications. Examples of such heterocyclic group and reference publications are recited in the following.

- 5-oxo- $\Delta^2$ -1,2,4-oxadiazolin-3-yl (or 2,5-dihydro-5-oxo-4H-1,2,4-oxadiazol-3-yl), 5-oxo- $\Delta^2$ -1,2,4-thiadiazolin-3-yl (or 2,5-dihydro-5-oxo-4H-1,2,4-thiadiazol-3-yl), 2-oxo- $\Delta^3$ -1,2,3,5-oxathiadiazolin-4-yl (or 2-oxo- $\Delta^3$ -1,2,4-oxathiadiazol-4-yl): Journal of Medicinal Chemistry, 39(26), 5228-35, 1996,
- 5-oxo- $\Delta^2$ -1,2,4-triazolin-3-yl: J Org Chem, 61(24), 8397-8401, 1996,
- 1-oxo- $\Delta^3$ -1,2,3,5-thiatriazolin-4-yl: Liebigs Ann Chem, 1376, 1980,
- 3-oxo- $\Delta^4$ -1,2,4-oxadiazolin-5-yl: EP145095,
- 5-oxo- $\Delta^2$ -1,3,4-oxadiazolin-2-yl: J Org Chem, 20, 412, 1955,
- 5-oxo- $\Delta^3$ -1,2,4-dioxazolin-3-yl: J Prakt Chem, 314, 145, 1972,
- 3-oxo- $\Delta^4$ -1,2,4-thiadiazolin-5-yl: JP-A-61-275271,
- 5-oxo- $\Delta^3$ -1,2,4-dithiazolin-3-yl: J Org Chem, 61(19), 6639-6645, 1996,
- 2-oxo- $\Delta^4$ -1,3,4-dioxazolin-5-yl: J Org Chem, 39, 2472, 1974,
- 2-oxo- $\Delta^4$ -1,3,4-oxathiazolin-5-yl: J Med Chem, 35(20), 3691-98, 1992,
- 5-oxo- $\Delta^2$ -1,3,4-thiadiazolin-2-yl: J Prakt Chem, 332(1), 55, 1990,
- 5-oxo- $\Delta^2$ -1,4,2-oxathiazolin-3-yl: J Org Chem, 31, 2417, 1966,

2-oxo- $\Delta^4$ -1,3,4-dithiazolin-5-yl: Tetrahedron Lett, 23, 5453, 1982,  
2-oxo- $\Delta^4$ -1,3,2,4-dioxathiazolin-5-yl: Tetrahedron Lett, 319, 1968,  
3,5-dioxoisooxazolidin-4-yl: Helv-Chim Acta, 1973, 48, 1965,  
2,5-dioxoimidazolidin-4-yl: Heterocycles, 43(1), 49-52, 1996,  
5 5-oxo-2-thioxoimidazolidin-4-yl: Heterocycles, 5, 391, 1983,  
2,4-dioxooxazolidin-5-yl: J Am Chem Soc, 73, 4752, 1951,  
4-oxo-2-thioxooxazolidin-5-yl: Chem Ber, 91, 300, 1958,  
2,4-dioxothiazolidin-5-yl: JP-A-57-123175,  
4-oxo-2-thioxothiazolidin-5-yl: Chem Pharm Bull, 30, 3563, 1982,

10 The Production Methods shown in the above-mentioned  
Production Methods 2 to 4 can be used for the synthesis of  
compounds other than benzimidazole of the formulas [I] and [II],  
such as compounds [II-2-15] and [II-2-16].

The compounds of the formulas [I], [II] and [III], 4-(4-  
15 fluorophenyl)-5-hydroxymethyl-2-methylthiazole and 4-(4-  
fluorophenyl)-5-chloromethyl-2-methylthiazole and production  
methods thereof of the present invention are explained in detail  
in the following by way of Examples. It is needless to say that  
the present invention is not limited by these Examples.

20 **Example 1**

Production of ethyl 2-[4-(3-bromophenoxy)phenyl]-1-  
cyclohexylbenzimidazole-5-carboxylate

**Step 1:** Production of ethyl 4-chloro-3-nitrobenzoate

4-Chloro-3-nitrobenzoic acid (300 g) was dissolved in ethyl  
25 alcohol (1500 ml) and concentrated sulfuric acid (100 ml) was  
added with ice-cooling. The mixture was refluxed under heating  
for 7 hr. The reaction mixture was poured into ice-cold water and  
the precipitated crystals were collected by filtration to give  
the title compound (332 g, yield 97%).

30  $^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 8.50(1H, d,  $J=2.1\text{Hz}$ ), 8.16(1H, dd,  $J=8.4$ ,  
2.1Hz), 7.63(1H, d,  $J=8.4\text{Hz}$ ), 4.43(2H, q,  $J=7.5\text{Hz}$ ), 1.42(3H, t,  
 $J=7.5\text{Hz}$ )

**Step 2:** Production of ethyl 4-cyclohexylamino-3-nitrobenzoate

Ethyl 4-chloro-3-nitrobenzoate (330 g) obtained in the  
35 previous step was dissolved in acetonitrile (1500 ml), and  
cyclohexylamine (220 g) and triethylamine (195 g) were added. The  
mixture was refluxed under heating overnight. The reaction  
mixture was poured into ice-cold water and the precipitated

crystals were collected by filtration to give the title compound (400 g, yield 94%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.87(1H, d, J=2.1Hz), 8.35-8.46(1H, m), 8.02(1H, dd, J=9.1, 2.1Hz), 6.87(1H, d, J=9.1Hz), 4.35(2H, q, J=7.1Hz), 3.65-3.50(1H, m), 2.14-1.29(10H, m), 1.38(3H, t, J=7.1Hz)

**Step 3: Production of ethyl 3-amino-4-cyclohexylaminobenzoate**

Ethyl 4-cyclohexylamino-3-nitrobenzoate (400 g) obtained in the previous step was dissolved in ethyl acetate (1500 ml) and ethyl alcohol (500 ml), and 7.5% palladium carbon (50% wet, 40 g) was added. The mixture was hydrogenated for 7 hr at atmospheric pressure. The catalyst was filtered off and the filtrate was concentrated under reduced pressure. Diisopropyl ether was added to the residue and the precipitated crystals were collected by filtration to give the title compound (289 g, yield 80%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.57(1H, dd, J=8.4, 1.9Hz), 7.41(1H, d, J=1.9Hz), 6.59(1H, d, J=8.4Hz), 4.30(2H, q, J=7.1Hz), 3.40-3.30(1H, m), 2.18-2.02(2H, m), 1.88-1.15(8H, m), 1.35(3H, t, J=7.1Hz)

**Step 4: Production of ethyl 3-[4-(3-bromophenoxy)benzoyl]amino-4-cyclohexylaminobenzoate**

4-(3-Bromophenoxy)benzoic acid (74 g) was dissolved in chloroform (500 ml), and oxalyl chloride (33 ml) and dimethylformamide (catalytic amount) were added. The mixture was stirred for 4 hr at room temperature. The reaction mixture was concentrated under reduced pressure and dissolved in dichloromethane (150 ml). The resulting solution was added dropwise to a solution of ethyl 3-amino-4-cyclohexylaminobenzoate (66 g) obtained in the previous step in dichloromethane (500 ml) and triethylamine (71 ml), and the mixture was stirred for 1 hr at room temperature. The reaction mixture was poured into water and extracted with dichloromethane. The organic layer was washed with saturated brine, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. Diethyl ether was added to the residue for crystallization and the crystals were collected by filtration to give the title compound (129 g, yield 95%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.00-7.78(4H, m), 7.66(1H, brs), 7.37-7.18(3H, m), 7.13-6.59(3H, m), 6.72(1H, d, J=8.7Hz), 4.50(1H,

brs), 4.29 (2H, q, J=7.2Hz), 3.36 (1H, m), 2.12-1.96 (2H, m), 1, 83-1.56 (3H, m), 1.47-1.12 (5H, m), 1.37 (3H, t, J=7.2Hz)

**Step 5: Production of ethyl 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate**

5 Ethyl 3-[4-(3-bromophenoxy)benzoyl]amino-4-cyclohexylaminobenzoate (129 g) obtained in the previous step was suspended in acetic acid (600 ml) and the resulting suspension was refluxed under heating for 3 hr. The reaction mixture was concentrated under reduced pressure. Water was added to the  
10 residue and the precipitated crystals were collected by filtration to give the title compound (124 g, yield 99%).  
<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.51 (1H, d, J=1.5Hz), 8.00 (1H, dd, J=8.4, 1.5Hz), 7.67 (1H, d, J=8.4Hz), 7.63 (2H, d, J=8.7Hz), 7.35-7.21 (3H, m), 7.17 (2H, d, J=8.7Hz), 7.14 (1H, m), 4.42 (2H, q, J=7.2Hz),  
15 4.38 (1H, m), 2.43-2.22 (2H, m), 2.07-1.87 (4H, m), 1.80 (1H, m), 1.42 (3H, t, J=7.2Hz), 1.40-1.27 (3H, m)

**Example 2**

Production of 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid

20 Ethyl 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (1.0 g) obtained in Example 1 was dissolved in tetrahydrofuran (10 ml) and ethyl alcohol (10 ml), and 4N sodium hydroxide (10 ml) was added. The mixture was refluxed under heating for 1 hr. The reaction mixture was  
25 concentrated under reduced pressure and water was added to the residue. The mixture was acidified with 6N hydrochloric acid and the precipitated crystals were collected by filtration to give the title compound (0.9 g, yield 96%).

melting point: 255-256°C

30 FAB-MS: 491 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): (12.75 (1H, brs), 8.24 (1H, s), 7.96 (1H, d, J=8.7Hz), 7.86 (1H, d, J=8.7Hz), 7.71 (2H, d, J=8.6Hz), 7.47-7.34 (3H, m), 7.24 (2H, d, J=8.6Hz), 7.20 (1H, m), 4.31 (1H, m), 2.38-2.18 (2H, m), 2.02-1.79 (4H, m), 1.65 (1H, m), 1.44-1.20 (3H, m)

35 **Example 3**

Production of ethyl 1-cyclohexyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylate

Ethyl 3-amino-4-cyclohexylaminobenzoate (130 g) obtained in Example 1, Step 3, and methyl 4-hydroxybenzimidate hydrochloride (139 g) were added to methyl alcohol (1500 ml), and the mixture was refluxed under heating for 4 hr. The reaction mixture was  
5 allowed to cool and the precipitated crystals were collected by filtration to give the title compound (131 g, yield 72%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 10.02(1H, brs), 8.21(1H, d, J=1.4Hz), 7.93(1H, d, J=8.6Hz), 7.83(1H, dd, J=8.6, 1.4Hz), 7.48(2H, d, J=8.6Hz), 6.95(2H, d, J=8.6Hz), 4.39-4.25(1H, m), 4.33(1H, q, J=7.0Hz),  
10 2.35-2.18(2H, m), 1.98-1.79(4H, m), 1.70-1.60(1H, m), 1.46-1.19(3H, m), 1.35(3H, t, J=7.0Hz)

#### Example 4

Production of ethyl 2-[4-(2-bromo-5-chlorobenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

15 2-Bromo-5-chlorobenzyl bromide prepared from 2-bromo-5-chlorotoluene (50 g), N-bromosuccinimide and N,N'-azobisisobutyronitrile, and ethyl 1-cyclohexyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylate (50 g) obtained in Example 3 were suspended in dimethylformamide (300 ml). Potassium  
20 carbonate (38 g) was added and the mixture was stirred for 1 hr at 80°C with heating. The reaction mixture was allowed to cool and then added to a mixed solvent of water-ethyl acetate. The precipitated crystals were collected by filtration to give the title compound (50 g, yield 64%).

25 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.50(1H, d, J=1.4Hz), 7.97(1H, dd, J=8.6, 1.4Hz), 7.70-7.57(5H, m), 7.20(1H, dd, J=8.4, 2.5Hz), 7.14(2H, d, J=8.7Hz), 5.17(2H, s), 4.46-4.30(1H, m), 4.41(2H, q, J=7.1Hz), 2.40-2.20(2H, m), 2.02-1.21(8H, m), 1.42(3H, t, J=7.1Hz)

#### Example 5

30 Production of ethyl 2-[4-[2-(4-chlorophenyl)-5-chlorobenzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Ethyl 2-[4-(2-bromo-5-chlorobenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (49 g) obtained in Example 4, 4-chlorophenylboronic acid (18 g) and tetrakis-  
35 (triphenylphosphine)palladium (10 g) were suspended in 1,2-dimethoxyethane (600 ml). Saturated aqueous sodium hydrogencarbonate solution (300 ml) was added and the mixture was refluxed under heating for 2 hr. Chloroform was added to the



reaction mixture. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate solution, water and saturated brine, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, chloroform:ethyl acetate = 97:3). Ethyl acetate and diisopropyl ether were added to the resulting oil for crystallization and the resulting crystals were collected by filtration to give the title compound (44 g, yield 85%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.49(1H, d, J=1.4Hz), 7.97(1H, dd, J=8.6, 1.6Hz), 7.70-7.60(2H, m), 7.55(2H, d, J=8.7Hz), 4.95(2H, s), 4.48-4.28(1H, m), 4.40(2H, m), 2.02-1.20(8H, m), 1.41(3H, t, J=7.1Hz)

#### Example 6

Production of 2-{4-[2-(4-chlorophenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid

Ethyl 2-{4-[2-(4-chlorophenyl)-5-chlorobenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (43 g) obtained in Example 5 was treated in the same manner as in Example 2 to give the title compound (33 g, yield 76%).

melting point: 243-244°C

FAB-MS: 571(MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.32(1H, s), 8.28(1H, d, J=8.9Hz), 8.05(1H, d, J=8.8Hz), 7.76-7.72(3H, m), 7.58-7.46(5H, m), 7.40(1H, d, J=8.3Hz), 7.24(2H, d, J=8.9Hz), 5.11(2H, s), 4.36(1H, m), 2.40-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-1.15(3H, m)

#### Example 7

Production of ethyl 2-[4-(2-bromo-5-methoxybenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Ethyl 1-cyclohexyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylate obtained in Example 3 and 2-bromo-5-methoxybenzyl bromide were treated in the same manner as in Example 4 to give the title compound (59 g).

#### Example 8

Production of ethyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate

Ethyl 2-[4-(2-bromo-5-methoxybenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate obtained in Example 7 was treated in the same manner as in Example 5 to give the title compound (48 g, yield 77%).

5 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.49(1H, d, J=1.4Hz), 7.97(1H, dd, J=8.6, 1.4Hz), 7.64(1H, d, J=8.6Hz), 7.54(2H, d, J=8.7Hz), 7.37(2H, d, J=8.6Hz), 7.31(2H, d, J=8.6Hz), 7.25(1H, d, J=8.4Hz), 7.19(1H, d, J=2.7Hz), 7.00(2H, d, J=8.7Hz), 6.97(1H, dd, J=8.4, 2.7Hz),  
10 4.98(2H, s), 4.41(2H, q, J=7.1Hz), 4.42-4.29(1H, m), 3.88(3H, s), 2.40-2.20(2H, m), 2.01-1.88(4H, m), 1.83-1.73(1H, m), 1.42(3H, t, J=7.1Hz), 1.41-1.25(3H, m)

#### Example 9

Production of 2-[4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid

15 Ethyl 2-[4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (52 g) obtained in Example 8 was treated in the same manner as in Example 2 to give the title compound (44 g, yield 89%).

melting point: 248-249°C

20 FAB-MS: 568 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.20(1H, s), 7.88(1H, d, J=8.7Hz),  
7.85(1H, d, J=8.7Hz), 7.57(d, 2H, J=8.6Hz), 7.46(2H, d, J=8.6Hz),  
7.44(2H, d, J=8.6Hz), 7.29(1H, d, J=8.5Hz), 7.24(1H, d, J=2.6Hz),  
7.11(2H, d, J=8.6Hz), 7.06(1H, dd, J=8.5, 2.6Hz), 5.04(2H, s),  
25 4.26(1H, m), 3.83(3H, s), 2.38-2.29(2H, m)

#### Example 10

Production of ethyl 1-cyclohexyl-2-[4-[(E)-2-phenylvinyl]phenyl]-benzimidazole-5-carboxylate

Ethyl 3-amino-4-cyclohexylaminobenzoate (500 mg) obtained in  
30 Example 1, Step 3, was dissolved in methyl alcohol (6 ml) and trans-4-stilbenecarbaldehyde (397 mg) was added under ice-cooling. The mixture was stirred overnight at room temperature. The reaction mixture was ice-cooled and benzofuroxan (259 mg) dissolved in acetonitrile (2 ml) was added. The mixture was  
35 stirred for 7 hr at 50°C. The reaction mixture was ice-cooled. After 1N sodium hydroxide was added, ethyl acetate was added and the mixture was extracted. The organic layer was washed with water and saturated brine, dried over anhydrous magnesium sulfate,

and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 4:1) to give the title compound (540 mg, yield 63%).

- 5 <sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.28(1H, d, J=1.4Hz), 8.01(1H, d, J=8.7Hz), 7.90-7.80(3H, m), 7.75-7.65(4H, m), 7.50-7.25(5H, m), 4.35(2H, q, J=7.0Hz), 4.31(1H, m), 2.40-2.20(2H, m), 2.00-1.80(4H, m), 1.63(1H, m), 1.40-1.20(3H, m), 1.36(3H, t, J=7.0Hz)

#### Example 11

- 10 Production of 1-cyclohexyl-2-{4-[(E)-2-phenylvinyl]phenyl}-benzimidazole-5-carboxylic acid

Ethyl 1-cyclohexyl-2-{4-[(E)-2-phenylvinyl]phenyl}-benzimidazole-5-carboxylate (127 mg) obtained in Example 10 was treated in the same manner as in Example 2 to give the title  
15 compound (116 mg, yield 97%).

melting point: not lower than 300°C

FAB-MS: 423 (MH<sup>+</sup>)

- <sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.25(1H, s), 7.96-7.29(13H, m), 4.33(1H, brt), 2.41-2.23(2H, m), 2.03-1.78(4H, m), 1.71-1.59(1H, m), 1.49-  
20 1.20(3H, m)

#### Example 12

Production of 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxylic acid

- In the same manner as in Examples 1 and 2, the title  
25 compound (700 mg) was obtained.

FAB-MS: 413 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.60(1H, s), 8.04(1H, d, J=9.0Hz), 7.63(2H, d, J=8.4Hz), 7.51-7.32(6H, m), 7.14(2H, d, J=9.0Hz), 5.16(2H, s), 5.03-4.89(1H, m), 2.41-1.63(8H, m)

#### 30 Example 13

Production of 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxamide

- 2-(4-Benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxylic acid (700 mg) obtained in Example 12 was dissolved in  
35 dimethylformamide (10 ml), and ammonium chloride (108 mg), 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (390 mg), 1-hydroxybenzotriazole (275 mg) and triethylamine (0.3 ml) were added. The mixture was stirred overnight at room temperature.

Water was added to the reaction mixture and the mixture was extracted with ethyl acetate. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, water and saturated brine, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. Ethyl acetate and diisopropyl ether were added to the residue for crystallization and the crystals were collected by filtration to give the title compound (571 mg, yield 81%).

melting point: 232-233°C

10 FAB-MS: 412 (MH+)

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.23 (1H, d, =1.5Hz), 7.86 (1H, dd, J=8.5, 1.5Hz), 7.65-7.30 (8H, m), 7.13 (2H, d, J=8.8Hz), 5.16 (2H, s), 4.93 (1H, quint, J=8.8Hz), 2.40-1.60 (8H, m)

#### Example 14

15 Production of 2-(4-benzyloxyphenyl)-5-cyano-1-cyclopentylbenzimidazole

In the same manner as in Example 1, the title compound (400 mg) was obtained.

FAB-MS: 394 (MH+)

20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.11 (1H, s), 7.68-7.30 (9H, m), 7.13 (2H, s), 5.16 (2H, s), 4.94 (1H, quint, J=8.9Hz), 2.35-1.60 (8H, m)

#### Example 15

Production of 2-(4-benzyloxyphenyl)-1-cyclopentylbenzimidazole-5-carboxamide oxime

25 2-(4-Benzyloxyphenyl)-5-cyano-1-cyclopentylbenzimidazole (400 mg) obtained in Example 14 was suspended in ethyl alcohol (3 ml) and water (1.5 ml), and hydroxylamine hydrochloride (141 mg) and sodium hydrogencarbonate (170 mg) were added. The mixture was refluxed under heating overnight. The reaction mixture was allowed to cool and the precipitated crystals were collected by filtration to give the title compound (312 mg, yield 71%).

melting point: 225-226°C

FAB-MS: 456 (MH+)

35 <sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.20 (1H, s), 7.50-7.31 (9H, m), 7.12 (2H, d, J=8.7Hz), 5.15 (2H, s), 4.94 (1H, quint, J=8.7Hz), 3.61 (3H, s), 3.40 (3H, s), 2.41-1.42 (8H, m)

#### Example 16

Production of ethyl 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylate

**Step 1:** Production of 4-(4-fluorophenyl)-5-hydroxymethyl-2-methylthiazole

5 Ethyl 4-(4-fluorophenyl)-2-methyl-5-thiazolecarboxylate (59 g) prepared by a known method (Chem. Pharm. Bull., 43(6), 947, 1995) was dissolved in tetrahydrofuran (700 ml). Lithium aluminum hydride (13 g) was added under ice-cooling and the mixture was stirred for 30 min. Water (13 ml), 15% sodium hydroxide (13 ml)  
10 and water (39 ml) were added successively to the reaction mixture, and the precipitated insoluble materials were filtered off. The filtrate was concentrated under reduced pressure to give the title compound (37 g, yield 71%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.60 (2H, dd, J=8.7, 6.6Hz), 7.11 (2H, t, J=8.7Hz), 4.80 (2H, s), 2.70 (3H, s)  
15

**Step 2:** Production of 5-chloromethyl-4-(4-fluorophenyl)-2-methylthiazole

4-(4-Fluorophenyl)-5-hydroxymethyl-2-methylthiazole (37 g) obtained in the previous step was dissolved in chloroform (500  
20 ml), and thionyl chloride (24 ml) and pyridine (2 ml) were added. The mixture was stirred for 3 hr at room temperature. The reaction mixture was poured into ice-cold water. The mixture was extracted with chloroform, and washed with water and saturated brine. The organic layer was dried over sodium sulfate, and  
25 concentrated under reduced pressure to give the title compound (29 g, yield 76%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.67 (2H, dd, J=8.8, 5.4Hz), 7.16 (2H, t, J=8.7Hz), 4.79 (2H, s), 2.73 (3H, s)

**Step 3:** Production of ethyl 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolyl]methoxy}phenyl}benzimidazole-5-carboxylate  
30

5-Chloromethyl-4-(4-fluorophenyl)-2-methylthiazole (28 g) obtained in the previous step and ethyl 1-cyclohexyl-2-(4-hydroxyphenyl)benzimidazole-5-carboxylate (36 g) obtained in  
35 Example 3 were treated in the same manner as in Example 4 to give the title compound (61 g, yield 100%).

APCI-MS: 570 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.25(1H, d, J=1.5Hz), 7.97(1H, d, J=8.7Hz), 7.86(1H, dd, J=8.6, 1.6Hz), 7.74(2H, dd, J=8.8, 5.5Hz), 7.62(2H, d, J=8.7Hz), 7.33(2H, t, J=8.9Hz), 7.22(2H, t, J=8.9Hz), 5.41(2H, s), 4.34(2H, q, J=7.1Hz), 4.31(1H, m), 2.71(3H, s),  
5 2.40-2.15(2H, m), 2.05-1.75(4H, m), 1.55-1.15(3H, m), 1.36(3H, t, J=7.1Hz)

#### Example 17

Production of 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolylmethoxy]phenyl}benzimidazole-5-carboxylic acid

10 Ethyl 1-cyclohexyl-2-{4-[4-(4-fluorophenyl)-2-methyl-5-thiazolylmethoxy]phenyl}benzimidazole-5-carboxylate (60 g) obtained in Example 16 was treated in the same manner as in Example 2 to give the title compound (39g, yield 69%).

melting point: 196-198°C

15 FAB-MS: 542 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 13.1(1H, brs), 8.34(1H, s), 8.29(1H, d, J=8.8Hz), 8.06(1H, d, J=8.7Hz), 7.80-7.72(4H, m), 7.36-7.31(4H, m), 5.46(2H, s), 4.38(1H, m), 2.72(3H, s), 2.45-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-  
20 1.20(3H, m)

#### Example 18

Production of ethyl 1-cyclohexyl-2-(2-fluoro-4-hydroxyphenyl)-benzimidazole-5-carboxylate

In the same manner as in Example 3, the title compound (50  
25 g) was obtained.

#### Example 19

Production of ethyl 2-{4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate

**Step 1** : Production of 3,3'-difluorobenzhydrol

30 To a stirred solution of magnesium strip (35.4 g) in THF (200 ml), iodine strip was added and the mixture was heated with stirring under nitrogen stream until most of color of iodine was disappeared. A solution of 3-fluoro-bromobenzene (250.0 g) in THF (1000 ml) was added dropwise over 2.5 hr while the temperature of  
35 the solution was maintained at 60°C. After the completion of the addition of the solution, the resulting mixture was refluxed for 1 hr with heating. The resulting Grignard solution was ice-cooled and a solution of ethyl formate (63.2 g) in THF (200 ml) was

added dropwise over 1 hr. After a stirring of the reaction solution for an additional 30 min, saturated aqueous ammonium chloride solution (700 ml) was added dropwise with ice-cooling and water (300 ml) was added. The mixture was stirred for 10 min.  
5 The organic layer and water layer were separated. Water layer was extracted with ethyl acetate, and the combined organic layer was washed with 2N hydrochloric acid, saturated aqueous sodium hydrogencarbonate and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, filtered, and the solvent  
10 was evaporated off under reduced pressure to give the title compound (156.2 g, yield 99%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.31(2H, td, J=7.9, 5.8Hz), 7.15-7.80(4H, m), 6.97-6.94(2H, m), 5.82(1H, d, J=3.3Hz), 2.30(1H, d, J=3.3Hz)

**Step 2: Production of 3,3'-difluorobenzhydryl chloride**

15 To a solution of 3,3'-difluorobenzhydrol (150.0 g) obtained in the previous step in toluene (400 ml), pyridine (539 mg) was added at room temperature. To the solution, thionyl chloride (89.1 g) was added dropwise over 1 hr at room temperature and the resulting solution was stirred for an additional 2 hr. The  
20 solution was heated so that the temperature of the solution was at 40°C, and then stirred for an additional 1.5 hr. Thionyl chloride (8.1 g) was added again and the mixture was stirred for 30 min. To the reaction mixture, water was added. The organic layer was separated, and washed with water, saturated aqueous  
25 sodium hydrogencarbonate and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, filtered, the solvent was evaporated off under reduced pressure to give the title compound (158.2 g, yield 97%).

30 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.32(2H, td, J=8.0, 5.9Hz), 7.18-7.10(4H, m), 7.01(2H, tdd, J=8.2, 2.5, 1.2Hz), 6.05(1H, s)

**Step 3: Production of ethyl 2-{4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate**

Ethyl 1-cyclohexyl-2-(2-fluoro-4-hydroxyphenyl)-benzimidazole-5-carboxylate (50 g) obtained in Example 18 and  
35 3,3'-difluorobenzhydryl chloride (34 g) obtained in the previous step were treated in the same manner as in Example 4 to give the title compound (76 g, yield 99%).

FAB-Ms: 585 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.24(1H, d, J=1.4Hz), 7.98(1H, d, J=8.7Hz), 7.88(1H, d, J=8.7Hz), 7.56(1H, t, J=8.6Hz), 7.50-7.40(6H, m), 6.82(1H, s), 4.34(2H, q, J=7.1Hz), 3.95(1H, m), 2.20-2.10(2H, m), 1.90-1.80(4H, m), 1.6(1H, m), 1.35(3H, t, J=7.2Hz), 1.30-1.20(3H, m)

#### Example 20

Production of 2-{4-(bis[3-fluorophenyl]methoxy)-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid

Ethyl 2-{4-[bis(3-fluorophenyl)methoxy]-2-fluorophenyl}-1-cyclohexylbenzimidazole-5-carboxylate (75 g) obtained in Example 19 was treated in the same manner as in Example 2 to give the title compound (48 g, yield 62%).

melting point: 242-243°C

FAB-MS: 557 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.29(1H, s), 8.16(1H, d, J=8.8Hz), 7.99(1H, d, J=8.7Hz), 7.66(1H, t, J=8.7Hz), 7.51-7.40(6H, m), 7.30(1H, d, J=12.1Hz), 7.20-7.14(3H, m), 6.88(1H, s), 4.07(1H, m), 2.40-2.10(2H, m), 2.00-1.75(4H, m), 1.70-1.55(1H, m), 1.50-1.15(3H, m)

#### Example 21

Production of ethyl 1-cyclopentyl-2-(4-nitrophenyl)benzimidazole-5-carboxylate

In the same manner as in Example 1, the title compound (12 g) was obtained.

#### Example 22

Production of ethyl 2-(4-aminophenyl)-1-cyclopentylbenzimidazole-5-carboxylate

Ethyl 1-cyclopentyl-2-(4-nitrophenyl)benzimidazole-5-carboxylate (12 g) obtained in Example 21 was dissolved in tetrahydrofuran (200 ml) and ethyl alcohol (50 ml), 7.5% palladium carbon (50% wet, 1 g) was added. The mixture was hydrogenated for 1 hr at atmospheric pressure. The catalyst was filtered off and the filtrate was concentrated under reduced pressure. Tetrahydrofuran was added to the residue to allow crystallization and the crystals were collected by filtration to give the title compound (11 g, yield 98%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.49(1H, d, J=1.3Hz), 7.95(1H, dd, J=8.5, 1.3Hz), 7.50-7.40(3H, m), 6.79(2H, d, J=4.6Hz), 4.97(1H, quint,



J=8.9Hz), 4.40 (2H, q, J=7.1Hz), 3.74 (2H, brs), 2.40-1.60 (8H, m), 1.41 (3H, t, J=7.1Hz)

#### Example 23

Production of ethyl 2-(4-benzoylaminophenyl)-1-cyclopentylbenzimidazole-5-carboxylate

Ethyl 1-cyclopentyl-2-(4-aminophenyl)benzimidazole-5-carboxylate (300 mg) obtained in Example 22 was dissolved in pyridine (3 ml) and chloroform (3 ml), and benzoyl chloride (127 mg) was added. The mixture was stirred for 30 min at room temperature. The reaction mixture was concentrated under reduced pressure and water was added to the residue to allow crystallization. The crystals were collected by filtration to give the title compound (403 mg, yield 100%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.58 (1H, s), 8.00 (1H, d, J=9.0Hz), 7.84 (2H, d, J=7.5Hz), 7.60-7.40 (6H, m), 7.14 (2H, d, J=7.5Hz), 4.84 (1H, quint, J=8.7Hz), 4.41 (2H, q, J=7.5Hz), 2.20-1.30 (8H, m), 1.41 (3H, t, J=7.5Hz)

#### Example 24

Production of 2-(4-benzoylaminophenyl)-1-cyclopentylbenzimidazole-5-carboxylic acid

Ethyl 2-(4-benzoylaminophenyl)-1-cyclopentylbenzimidazole-5-carboxylate (200 mg) obtained in Example 23 was treated in the same manner as in Example 2 to give the title compound (131 mg, yield 70%).

melting point: not lower than 300°C

FAB-MS: 426 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 10.75 (1H, s), 8.35 (1H, s), 8.15 and 7.85 (4H, ABq, J=8.9Hz), 8.10-7.98 (4H, m), 7.70-7.55 (3H, m), 5.02 (1H, quint, J=8.7Hz), 2.36-2.15 (4H, m), 2.14-1.95 (2H, m), 1.80-1.62 (2H, m)

#### Example 25

Production of ethyl 2-{4-[3-(3-chlorophenyl)phenoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate

Ethyl 2-[4-(3-bromophenoxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (65 g) obtained in Example 1 and 3-chlorophenylboronic acid (23 g) were treated in the same manner as in Example 5 to give the title compound (59 g, yield 85%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.51(1H, d, J=1.8Hz), 7.99(1H, dd, J=8.7, 1.8Hz), 7.71-7.55(4H, m), 7.51-7.43(2H, m), 7.43-7.27(4H, m), 7.19(1H, d, J=8.4Hz), 7.12(1H, m), 4.41(2H, q, J=7.2Hz), 4.39(1H, m), 2.42-2.22(2H, m), 2.03-1.87(4H, m), 1.79(1H, m), 1.42(3H, t, J=7.2Hz), 1.39-1.29(3H, m)

#### Example 26

Production of 2-[4-[3-(3-chlorophenyl)phenoxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid

Ethyl 2-[4-[3-(3-chlorophenyl)phenoxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (59 g) obtained in Example 25 was treated in the same manner as in Example 2 to give the title compound (43 g, yield 76%).

melting point: 253-254°C

FAB-MS: 523 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 12.82(1H, brs), 8.24(1H, d, J=1.3Hz), 7.98(1H, d, J=8.7Hz), 7.89(1H, dd, J=8.7, 1.3Hz), 7.78(1H, s), 7.72(2H, d, J=9.7Hz), 7.70(1H, m), 7.64-7.42(5H, m), 7.25(2H, d, J=8.7Hz), 7.20(1H, m), 4.33(1H, m), 2.39-2.17(2H, m), 2.00-1.76(4H, m), 1.65(1H, m), 1.50-1.22(3H, m)

#### Example 27

Production of ethyl 2-[4-(3-acetoxyphenyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

In the same manner as in Example 1, the title compound (87 g) was obtained.

#### Example 28

Production of ethyl 1-cyclohexyl-2-[4-(3-hydroxyphenyloxy)-phenyl]benzimidazole-5-carboxylate

Ethyl 2-[4-(3-acetoxyphenyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (87 g) obtained in Example 27 was dissolved in methyl alcohol (250 ml) and tetrahydrofuran (250 ml), and potassium carbonate (31 g) was added. The mixture was stirred for 30 min at room temperature. The insoluble materials were filtered off and the filtrate was concentrated under reduced pressure. Water was added to the residue and the mixture was neutralized with 2N hydrochloric acid. The precipitated crystals were collected by filtration to give the title compound (78 g, yield 97%).

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 9.71(1H, s), 7.98(1H, d, J=8.7Hz), 7.87(1H, d, J=8.7Hz), 7.68(2H, d, J=8.6Hz), 7.24(1H, t, J=8.1Hz), 7.18(2H, d, J=8.6Hz), 6.63(1H, d, J=8.1Hz), 6.57(1H, d, J=8.1Hz), 6.51(1H, s), 4.38-4.23(1H, m), 4.35(2H, q, J=6.9Hz), 2.36-2.18(2H, m), 1.99-1.78(4H, m), 1.71-1.59(1H, m), 1.45-1.20(3H, m), 1.36(3H, t, J=6.9Hz)

#### Example 29

Production of ethyl 1-cyclohexyl-2-{4-[3-(4-pyridylmethoxy)phenyloxy]phenyl}benzimidazole-5-carboxylate

10 Ethyl 1-cyclohexyl-2-[4-(3-hydroxyphenyloxy)phenyl]-benzimidazole-5-carboxylate (78 g) obtained in Example 28 was suspended in dimethylformamide (800 ml), and sodium hydride (60% oil, 14 g) was added under ice-cooling. The mixture was stirred for 1 hr at room temperature. After the reaction mixture was ice-cooled, 4-chloromethylpyridine hydrochloride (29 g) was added and the mixture was stirred for 30 min. The mixture was then stirred overnight at room temperature. Water was added to the reaction mixture and the precipitated crystals were collected by filtration. The resulting crystals were recrystallized from ethyl alcohol to give the title compound (77 g, yield 82%).

20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.63(2H, d, J=6.0Hz), 8.51(1H, s), 7.99(1H, d, J=8.7Hz), 7.66(2H, d, J=8.7Hz), 7.62(2H, d, J=8.7Hz), 7.36(2H, d, J=8.7Hz), 7.31(1H, t, J=8.2Hz), 7.26(1H, s), 7.16(2H, d, J=8.7Hz), 6.79-6.70(3H, m), 5.09(2H, s), 4.47-4.31(1H, m), 4.42(2H, q, J=7.0Hz), 2.42-2.22(2H, m), 2.04-1.71(5H, m), 1.45-1.25(3H, m), 1.42(3H, t, J=7.0Hz)

#### Example 30

Production of 1-cyclohexyl-2-{4-[3-(4-pyridylmethoxy)phenyloxy]phenyl}benzimidazole-5-carboxylic acid

30 Ethyl 1-cyclohexyl-2-{4-[3-(4-pyridylmethoxy)phenyloxy]phenyl}benzimidazole-5-carboxylate (60 g) obtained in Example 29 was treated in the same manner as in Example 2 to give the title compound (54 g, yield 75%).

melting point: 235-237°C

35 FAB-MS: 520 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.58(2H, d, J=6.0Hz), 8.23(1H, s), 7.96 and 7.86(2H, ABq, J=8.7Hz), 7.68 and 7.17(4H, A'B'q, J=8.7Hz), 7.44(2H, d, J=8.7Hz), 7.39(1H, t, J=8.3Hz), 6.90(1H, d, J=8.1Hz),

6.84 (1H, s), 6.75 (1H, d, J=8.1Hz), 5.22 (2H, s), 4.40-4.22 (1H, m),  
2.40-2.19 (2H, m), 2.00-1.80 (4H, m)

#### **Example 241**

Production of methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate  
**Step 1:** Production of 2-bromo-5-methoxybenzaldehyde

3-Methoxybenzaldehyde (15 g) was dissolved in acetic acid (75 ml), and a solution of bromine (5.7 ml) dissolved in acetic acid (15 ml) was added dropwise. The mixture was stirred  
10 overnight at room temperature and water (150 ml) was added to the reaction mixture. The precipitated crystals were collected by filtration, washed with water and dried under reduced pressure to give the title compound (21 g, yield 88%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 10.31 (1H, s), 7.52 (1H, d, J=8.8Hz),  
15 7.41 (1H, d, J=3.3Hz), 7.03 (1H, dd, J=8.8, 3.3Hz), 3.48 (3H, s)

**Step 2:** Production of 2-(4-chlorophenyl)-5-methoxybenzaldehyde

2-Bromo-5-methoxybenzaldehyde (10 g) obtained in the previous step was treated in the same method as in Example 5 to give the title compound (11 g, yield 96%).

20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 9.92 (1H, s), 7.50 (1H, d, J=2.6Hz), 7.48-7.14 (6H, m), 3.90 (3H, s)

**Step 3:** Production of 2-(4-chlorophenyl)-5-methoxybenzyl alcohol

2-(4-Chlorophenyl)-5-methoxybenzaldehyde (10 g) obtained in the previous step was dissolved in tetrahydrofuran (30 ml). The  
25 solution was added dropwise to a suspension of sodium borohydride (620 mg) in isopropyl alcohol (50 ml) and the mixture was stirred for 1 hr. The solvent was evaporated under reduced pressure and water was added to the residue. The precipitated crystals were collected by filtration and dried under reduced pressure. The  
30 resulting crystals were recrystallized from a mixture of methanol and water to give the title compound (9.2 g, yield 91%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.37 (2H, d, J=8.6Hz), 7.27 (2H, d, J=8.6Hz),  
7.17 (1H, d, J=8.6Hz), 7.11 (1H, d, J=2.6Hz), 6.89 (1H, dd, J=8.6,  
2.6Hz), 4.57 (2H, s), 3.86 (3H, s)

35 **Step 4:** Production of 2-(4-chlorophenyl)-5-methoxybenzyl chloride

2-(4-Chlorophenyl)-5-methoxybenzyl alcohol (20 g) obtained in the previous step was dissolved in ethyl acetate (100 ml) and pyridine (0.5 ml), and thionyl chloride (11 ml) was added

dropwise. The mixture was stirred for 1 hr. Water was added to the reaction mixture and the mixture was extracted with ethyl acetate. The organic layer was washed with water, saturated aqueous sodium hydrogencarbonate, water and saturated brine, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. Isopropyl alcohol was added to the residue to allow crystallization. The resulting crystals were collected by filtration and dried under reduced pressure to give the title compound (16 g, yield 74%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.43-7.29(4H, m), 7.17(1H, d, J=8.6Hz), 7.05(1H, d, J=2.6Hz), 6.96-6.89(1H, m), 4.46(2H, s), 3.86(3H, s)

**Step 5:** Production of methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate  
2-(4-Chlorophenyl)-5-methoxybenzyl chloride (4.0 g) obtained in the previous step and methyl 1-cyclohexyl-2-(4-hydroxyphenyl)-benzimidazole-5-carboxylate (5.0 g) obtained in the same manner as in Example 3 were treated in the same manner as in Example 4 to give the title compound (6.0 g, yield 72%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.48(1H, s), 8.00-7.93(1H, m), 7.68-7.62(1H, m), 7.54(2H, d, J=9.0Hz), 7.41-7.16(6H, m), 7.04-6.93(3H, m), 4.97(2H, s), 4.36(1H, m), 3.94(3H, s), 3.87(3H, s), 2.39-2.21(2H, m), 2.02-1.88(4H, m), 1.85-1.45(4H, m)

#### Example 242

Production of 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride

Methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (5.0 g) obtained in Example 241 was treated in the same manner as in Example 2 to give the title compound (5.1 g, yield 98%).

APCI-MS: 568 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.30(1H, d, J=1.4Hz), 8.24(1H, d, J=8.7Hz), 8.03(1H, d, J=8.7Hz), 7.72(2H, d, J=8.7Hz), 7.51-7.39(4H, m), 7.34-7.18(4H, m), 7.11-7.03(1H, m), 5.08(2H, s), 4.35(1H, m), 3.83(3H, m), 2.40-2.18(2H, m), 2.10-1.96(2H, m), 1.93-1.78(2Hm), 1.72-1.18(4H, m)

#### Example 243

Production of ethyl 2-{4-[3-(4-chlorophenyl)pyridin-2-ylmethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate

**Step 1: Production of methyl 3-hydroxypicolinate**

3-Hydroxypicolinic acid (1.0 g) was suspended in methanol (10 ml) and concentrated sulfuric acid (1.0 ml) was added. The mixture was refluxed under heating for 5 hr. The reaction mixture was ice-cooled, neutralized with saturated aqueous sodium hydrogencarbonate, and extracted with chloroform. The organic layer was washed with water and saturated brine, and dried over anhydrous magnesium sulfate. The solvent was evaporated under reduced pressure to give the title compound (711 mg, yield 64%).  
10 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 10.63(1H, s), 8.28(1H, dd, J=3.7, 1.8Hz), 7.47-7.35(2H, m), 4.06(3H, s)

**Step 2: Production of methyl 3-(trifluoromethylsulfonyloxy)-pyridine-2-carboxylate**

Methyl 3-hydroxypicolinate (710 mg) obtained in the previous step and triethylamine (0.77 ml) were dissolved in dichloromethane (7 ml), and trifluoromethanesulfonic anhydride (0.86 ml) was added under ice-cooling. The reaction mixture was allowed to warm to room temperature and the mixture was stirred for 2 hr. Water was added to the reaction mixture and the mixture was extracted with ethyl acetate. The organic layer was washed with saturated brine and dried over anhydrous magnesium sulfate. The solvent was evaporated under reduced pressure to give the title compound (1.2 g, yield 90%).  
20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.80-8.73(1H, m), 7.75-7.70(1H, m), 7.63(1H, dd, J=8.2, 4.5Hz), 4.05(3H, s)

**Step 3: Production of methyl 3-(4-chlorophenyl)pyridine-2-carboxylate**

Methyl 3-(trifluoromethylsulfonyloxy)pyridine-2-carboxylate (1.2 g) obtained in the previous step was treated in the same manner as in Example 5 to give the title compound (728 mg, yield 69%).  
30 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.73-8.66(1H, m), 7.77-7.68(1H, m), 7.49(1H, dd, J=7.8, 4.5Hz), 7.46-7.37(2H, m), 7.32-7.23(2H, m), 3.80(3H, s)

**Step 4: Production of [3-(4-chlorophenyl)pyridin-2-yl]methanol**

Methyl 3-(4-chlorophenyl)pyridine-2-carboxylate (720 mg) obtained in the previous step was dissolved in tetrahydrofuran (10 ml) and the solution was ice-cooled. Lithium aluminum hydride

(160 mg) was added to the solution and the mixture was stirred for 1 hr. To the reaction mixture were added successively water (1.6 ml), 15% sodium hydroxide (1.6 ml) and water (4.8 ml). The insoluble materials were filtered off and the filtrate was  
5 concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 1:1) to give the title compound (208 mg, yield 32%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.60 (1H, dd, J=4.8, 1.5Hz), 7.60-7.55 (1H,  
10 m), 7.40-7.48 (2H, m), 7.29-7.36 (1H, m), 7.27-7.20 (3H, m), 4.63 (2H, s)

**Step 5:** Production of ethyl 2-{4-[3-(4-chlorophenyl)pyridin-2-ylmethoxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate

[3-(4-Chlorophenyl)pyridin-2-yl]methanol (200 mg) obtained  
15 in the previous step was dissolved in chloroform (3 ml), and thionyl chloride (0.13 ml) and pyridine (catalytic amount) were added. The mixture was stirred for 1 hr at room temperature and concentrated under reduced pressure. The residue was dissolved in dimethylformamide (3 ml), and ethyl 1-cyclohexyl-2-(4-  
20 hydroxyphenyl)benzimidazole-5-carboxylate (232 mg) obtained in the same manner as in Example 3 and potassium carbonate (250 mg) were added. The mixture was stirred for 3 hr with heating at 80°C. The reaction mixture was then allowed to cool. Water was added and the mixture was extracted with ethyl acetate. The organic  
25 layer was washed with water and saturated brine, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 1:2) to give the title compound (246 mg, yield 68%).

30 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.71 (1H, dd, J=4.7, 1.4Hz), 8.49 (1H, d, J=2.1Hz), 7.96 (1H, d, J=10.2Hz), 7.71-7.62 (2H, m), 7.53 (2H, d, J=8.7Hz), 7.45-7.34 (5H, m); 7.04 (2H, d, J=8.7Hz), 5.14 (2H, s), 4.48-4.29 (3H, m), 2.38-2.19 (2H, m), 2.02-1.22 (11H, m)

#### **Example 244**

35 Production of methyl 2-[4-(2-bromo-5-tert-butoxycarbonylbenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

**Step 1:** Production of tert-butyl 4-bromo-3-methylbenzoate

4-Bromo-3-methylbenzoic acid (25 g) was suspended in dichloromethane (200 ml), and oxalyl chloride (12 ml) and dimethylformamide (catalytic amount) were added. The mixture was stirred for 2 hr at room temperature and the solvent was  
5 evaporated under reduced pressure. The residue was dissolved in tetrahydrofuran (200 ml) and the solution was ice-cooled. To the solution was added dropwise a solution of potassium tert-butoxide dissolved in tetrahydrofuran (150 ml) and the mixture was stirred for 30 min. Water was added to the reaction mixture and the  
10 mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated brine, and dried over anhydrous magnesium sulfate. The solvent was evaporated under reduced pressure to give the title compound (27 g, yield 85%).

$^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 7.83(1H, d,  $J=2.2\text{Hz}$ ), 7.67-7.53(2H, m),  
15 2.43(3H, s), 1.58(9H, s)

**Step 2:** Production of methyl 2-[4-(2-bromo-5-tert-butoxycarbonylbenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

tert-Butyl 4-bromo-3-methylbenzoate (7.0 g) obtained in the  
20 previous step and methyl 1-cyclohexyl-2-(4-hydroxyphenyl)-benzimidazole-5-carboxylate (6.3 g) obtained in the same manner as in Example 3 were treated in the same manner as in Example 4 to give the title compound (8.8 g, yield 77%).

$^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 8.49(1H, d,  $J=1.5\text{Hz}$ ), 8.21(1H, d,  $J=2.1\text{Hz}$ ),  
25 7.97(1H, d,  $J=10.2\text{Hz}$ ), 7.82(1H, d,  $J=10.2\text{Hz}$ ), 7.71-7.58(4H, m), 7.16(2H, d,  $J=8.7\text{Hz}$ ), 5.23(2H, s), 4.38(1H, m), 3.95(3H, s), 2.40-2.23(2H, m), 2.04-1.90(4H, m), 1.84-1.73(1H, m), 1.59(9H, s), 1.44-1.27(3H, m)

#### Example 245

30 Production of methyl 2-[4-[5-tert-butoxycarbonyl-2-(4-chlorophenyl)benzyloxy]phenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Methyl 2-[4-(2-bromo-5-tert-butoxycarbonylbenzyloxy)phenyl]-1-cyclohexylbenzimidazole-5-carboxylate (4.5 g) obtained in  
35 Example 244 was treated in the same manner as in Example 5 to give the title compound (3.6 g, yield 76%).

$^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 8.48(1H, s), 8.27(1H, d,  $J=1.8\text{Hz}$ ), 8.04(1H, dd,  $J=7.9, 1.5\text{Hz}$ ), 7.96(1H, dd,  $J=7.0, 1.5\text{Hz}$ ), 7.65(1H, d,



J=8.6Hz), 7.55(2H, d, J=8.6Hz), 7.43-7.32(5H, m), 7.01(2H, d, J=8.6Hz), 4.99(2H, s), 4.43-4.29(1H, m), 3.95(3H, s), 2.41-2.21(2H, m), 2.02-1.89(4H, m), 1.82-1.73(1H, m), 1.62(9H, s), 1.46-1.28(3H, m)

#### 5 **Example 246**

Production of methyl 2-{4-[5-carboxy-2-(4-chlorophenyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate hydrochloride

Methyl 2-{4-[5-tert-butoxycarbonyl-2-(4-chlorophenyl)-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (3.5 g)  
10 obtained in Example 245 was dissolved in dichloromethane (35 ml), and trifluoroacetic acid (35 ml) was added. The mixture was stirred for 1 hr at room temperature and the reaction mixture was concentrated under reduced pressure. The residue was dissolved in  
15 ethyl acetate, and 4N hydrochloric acid-ethyl acetate was added. The precipitated crystals were collected by filtration and dried under reduced pressure to give the title compound (3.3 g, yield 97%).

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.33(1H, d, J=1.5Hz), 8.29(1H, s),  
20 8.24(1H, d, J=1.8Hz), 8.09-8.00(2H, m), 7.74(2H, d, J=8.6Hz), 7.61-7.44(5H, m), 7.24(2H, d, J=8.6Hz), 5.19(2H, s), 4.36(1H, m), 3.93(3H, s), 2.37-1.21(10H, m)

#### **Example 247**

Production of methyl 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoyl-benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate  
25

Methyl 2-{4-[5-carboxy-2-(4-chlorophenyl)benzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate hydrochloride (400 mg)  
obtained in Example 246 was suspended in dichloromethane (5 ml), and oxalyl chloride (0.08 ml) and dimethylformamide (catalytic  
30 amount) were added. The mixture was stirred for 2 hr at room temperature. The reaction mixture was concentrated under reduced pressure and the residue was dissolved in dichloromethane (5 ml). The resulting solution was added dropwise to a mixed solution of 40% aqueous methylamine solution (5 ml) and tetrahydrofuran (5  
35 ml) under ice-cooling. The reaction mixture was stirred for 1 hr and concentrated under reduced pressure. Water was added to the residue and the mixture was extracted with ethyl acetate. The organic layer was washed with water, saturated aqueous sodium

hydrogencarbonate and saturated brine, and dried over anhydrous magnesium sulfate. The solvent was evaporated under reduced pressure and the residue was crystallized from ethyl acetate and diisopropyl ether. The crystals were collected by filtration and  
5 dried under reduced pressure to give the title compound (335 mg, yield 86%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.47(1H, s), 8.06(1H, d, J=1.8Hz), 7.96(1H, dd, J=8.6, 1.5Hz), 7.82(1H, dd, J=8.2, 2.2Hz), 7.64(1H, d, J=8.6Hz), 7.54(2H, d, J=9.0Hz), 7.44-7.31(5H, m), 6.99(2H, d, J=9.0Hz),  
10 6.35-6.26(1H, m), 5.00(2H, s), 4.35(1H, m), 3.95(3H, s), 3.05(3H, d, J=4.8Hz), 2.40-1.24(10H, m)

#### Example 248

Production of 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate hydrochloride  
15

Methyl 2-{4-[2-(4-chlorophenyl)-5-methylcarbamoylbenzyloxy]phenyl}-1-cyclohexylbenzimidazole-5-carboxylate (150 mg) obtained in Example 247 and tetrahydrofuran (2 ml) were treated in the same manner as in Example 2 to give the title compound (141 mg,  
20 yield 90%).

APCI-MS: 594(MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.65-8.58(1H, m), 8.27(1H, d, J=1.5Hz), 8.21(1H, d, J=8.2Hz), 8.15(1H, d, J=1.5Hz), 8.05-7.90(2H, m), 7.70(2H, d, J=8.6Hz), 7.56-7.43(5H, m), 7.21(2H, d, J=8.6Hz),  
25 5.14(2H, s), 4.34(1H, m), 2.81(3H, d, J=4.5Hz), 2.39-1.19(10H, m)

#### Example 336

Production of methyl 2-[4-(2-bromo-5-nitrobenzyloxy)-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Commercially available 2-bromo-5-nitrotoluene was  
30 dissolved in carbon tetrachloride (30 ml), and N-bromosuccinimide (2.9 g) and N,N'-azobisisobutyronitrile (228 mg) were added, which was followed by refluxing under heating overnight. The reaction mixture was allowed to cool, water was added and the mixture was extracted with chloroform. The organic layer was  
35 dried over magnesium sulfate and concentrated under reduced pressure. The residue was dissolved in dimethylformamide (30 ml) and methyl 2-(2-fluoro-4-hydroxyphenyl)-1-cyclohexylbenzimidazole-5-carboxylate (3.8 g) obtained in the

same manner as in Example 3 and potassium carbonate (3.8 g) were added, which was followed by stirring at 80°C for 1 hr. The reaction mixture was allowed to cool, water was added and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated brine, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (n-hexane:ethyl acetate = 1:1) to give the title compound (3.7 g, yield 61%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.55-8.45(2H, m), 8.15-8.05(1H, m), 7.99(1H, dd, J=8.6Hz, 1.5Hz), 7.70-7.55(2H, m), 7.05-6.85(2H, m), 5.24(2H, s), 4.06(1H, m), 3.95(3H, s), 2.35-2.15(2H, m), 2.05-1.85(4H, m), 1.80-1.70(1H, m), 1.45-1.20(3H, m)

#### Example 337

Production of methyl 2-[4-{2-(4-chlorophenyl)-5-nitrobenzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Methyl 2-[4-(2-bromo-5-nitrobenzyloxy)-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (2.0 g) obtained in Example 336, 4-chlorophenylboronic acid (590 mg) and tetrakis(triphenylphosphine)palladium (396 mg) were suspended in dimethoxyethane (40 ml), and saturated aqueous sodium hydrogencarbonate solution (20 ml) was added, which was followed by refluxing under heating for 1 hr. The reaction mixture was allowed to cool, water was added and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (n-hexane:ethyl acetate = 2:1) to give the title compound (1.9 g, yield 90%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.55(1H, d, J=2.3Hz), 8.49(1H, d, J=1.4Hz), 8.29(1H, dd, J=8.4Hz, 2.3Hz), 7.98(1H, dd, J=8.6Hz, 1.5Hz), 7.60-7.30(6H, m), 6.85-6.70(2H, m), 5.03(2H, s), 4.02(1H, m), 3.95(3H, s), 2.35-2.10(2H, m), 2.05-1.70(5H, m), 1.40-1.20(3H, m)

#### Example 338

Production of methyl 2-[4-{5-amino-2-(4-chlorophenyl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Methyl 2-[4-{2-(4-chlorophenyl)-5-nitrobenzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (1.9 g)

obtained in Example 337 was suspended in ethanol (40 ml), and tin(II) chloride dihydrate (3.5 g) was added, which was followed by refluxing under heating for 30 min. The reaction mixture was concentrated under reduced pressure, 4N sodium hydroxide was added and the mixture was extracted with chloroform. The organic layer was washed with 2N sodium hydroxide and water, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. Diisopropyl ether was added to the residue, and the precipitated crystals were collected by filtration to give the title compound (1.5 g, yield 82%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.49(1H, d, J=1.2Hz), 7.98(1H, dd, J=9.0, 1.5Hz), 7.66(1H, d, J=8.7Hz), 7.49(1H, t, J=8.4Hz), 7.40-7.20(3H, m), 7.13(1H, d, J=8.1Hz), 6.92(1H, d, J=2.7Hz), 6.85-6.65(4H, m), 4.92(2H, s), 4.03(1H, m), 3.95(3H, s), 3.82(2H, brs), 2.30-2.10(2H, m), 2.05-1.80(4H, m), 1.80-1.70(1H, m), 1.40-1.10(3H, m)

#### Example 339

Production of methyl 2-[4-{2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate

Methyl 2-[4-{5-amino-2-(4-chlorophenyl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (500 mg) obtained in Example 338 and triethylamine (0.14 ml) were dissolved in chloroform (5 ml), and commercially available chlorobutyl chloride (0.1 ml) was added under ice-cooling, which was followed by stirring at room temperature for 3 hr. Water was added to the reaction mixture and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated brine, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was dissolved in dimethylformamide (6 ml) and potassium carbonate (244 mg) was added, which was followed by stirring at 80°C for 1 hr. The reaction mixture was allowed to cool, water was added and the precipitated crystals were collected by filtration to give the title compound (502 mg, yield 89%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 4.89(1H, d, J=1.5Hz), 7.98(1H, dd, J=8.6Hz, 1.6Hz), 7.72(1H, d, J=2.2Hz), 7.75-7.65(2H, m), 7.49(1H, t, J=8.3Hz), 7.45-7.20(5H, m), 6.85-7.65(2H, m), 4.99(2H, s), 4.10-

3.85 (6H, m), 2.66 (2H, t, J=7.8Hz), 2.30-2.15 (4H, m), 2.00-1.85 (4H, m), 1.80-1.70 (1H, m), 1.45-1.20 (3H, m)

#### Example 340

Production of 2-[4-{2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylic acid hydrochloride

Methyl 2-[4-{2-(4-chlorophenyl)-5-(2-oxopyrrolidin-1-yl)benzyloxy}-2-fluorophenyl]-1-cyclohexylbenzimidazole-5-carboxylate (200 mg) obtained in Example 339 was treated in the same manner as in Example 2 to give the title compound (182 mg, yield 87%).

Ms: 638 (M+1)

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.28 (1H, d, J=1.3Hz), 8.10 (1H, d, J=8.7Hz), 8.05-7.90 (2H, m), 7.77 (1H, dd, J=8.4Hz, 2.2Hz), 7.61 (1H, t, J=8.5Hz), 7.55-7.35 (5H, m), 7.00-7.20 (2H, m), 5.09 (2H, s), 4.06 (1H, m), 3.90 (2H, t, J=6.9Hz), 2.60-2.45 (2H, m), 2.30-2.00 (4H, m), 1.95-1.75 (4H, m), 1.70-1.55 (1H, m), 1.45-1.15 (3H, m)

In the same manner as in Examples 1-30, 241-248 and 336-340 and optionally using other conventional methods, where necessary, the compounds of Examples 31-240, 249-335, 341-446, 701-703 and 1001-1559 were obtained. The chemical structures and properties are shown in Table 1 to 177, 185 to 212, 219 to 221 and 225 to 260.

#### Example 501

Production of methyl 2-[4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl]-1-cyclohexyl-1H-indole-5-carboxylate

**Step 1:** Production of methyl 3-bromo-4-cyclohexylaminobenzoate

3-Bromo-4-fluorobenzoic acid (2.0 g) was dissolved in methanol (20 ml) and concentrated sulfuric acid (2 ml) was added.

The mixture was refluxed for 3 hr. The reaction mixture was poured into ice-cold water and extracted with ethyl acetate (50 ml). The organic layer was washed with water (30 ml) and saturated brine (30 ml), and dried over sodium sulfate. After filtration, the solvent was evaporated under reduced pressure.

The residue was dissolved in dimethyl sulfoxide (20 ml) and cyclohexylamine (10.3 ml) was added. The mixture was stirred overnight at 120°C. The reaction mixture was poured into 10% aqueous citric acid solution (100 ml) and extracted with ethyl

acetate (100 ml). The organic layer was washed with water (50 ml) and saturated brine (50 ml), and dried over sodium sulfate. After filtration, the solvent was evaporated under reduced pressure and the residue was purified by silica gel flash chromatography

5 (developing solvent, n-hexane:ethyl acetate = 10:1) to give the title compound (2.6 g, yield 92%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.10(1H, d, J=1.9Hz), 7.83(1H, dd, J=1.9Hz, 8.6Hz), 6.59(1H, d, J=8.7Hz), 4.73(1H, brd, J=7.3Hz), 3.85(3H, s), 3.38(1H, m), 2.10-2.00(2H, m), 1.90-1.20(8H, m)

10 **Step 2:** Production of 4'-chloro-2-(4-iodophenoxymethyl)-4-methoxybiphenyl

4-Iodophenol (5.0 g) was dissolved in acetone (50 ml), and potassium carbonate (4.7 g) and 4'-chloro-2-chloromethyl-4-methoxybiphenyl (6.0 g) obtained in Example 241, Step 4 were  
15 added. The mixture was refluxed for 10 hr. The reaction mixture was concentrated and 4N aqueous sodium hydroxide solution (50 ml) was added. The precipitated crystals were collected by filtration, washed with water, and dried under reduced pressure to give the title compound (10.0 g, yield 98%).

20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.52(2H, d, J=8.9Hz), 7.35(2H, d, J=8.5Hz), 7.27-7.20(3H, m), 7.12(1H, s), 6.95(1H, d, J=8.5Hz), 6.62(2H, d, J=8.9Hz), 4.84(2H, s), 3.85(3H, s)

**Step 3:** Production of [4-(4'-chloro-4-methoxybiphenyl-2-ylmethoxy)phenylethynyl]trimethylsilane

25 4'-Chloro-2-(4-iodophenoxymethyl)-4-methoxybiphenyl (7.0 g) obtained in the previous step was dissolved in acetonitrile (50 ml), and trimethylsilylacetylene (2.3 g), tetrakis-(triphenylphosphine)palladium complex (1.8 g), copper(I) iodide (0.6 g) and triethylamine (50 ml) were added. The mixture was  
30 stirred overnight at room temperature and concentrated. Water (30 ml) was added and the mixture was extracted with ethyl acetate (50 ml). The organic layer was washed with water (30 ml) and saturated brine (30 ml) and dried over sodium sulfate. After filtration, the solvent was evaporated under reduced pressure and  
35 the residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 10:1) to give the title compound (5.1 g, yield 79%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.37(2H, d, J=8.9Hz), 7.34(2H, d, J=8.2Hz), 7.28-7.21(3H, m), 7.13(1H, s), 6.94(1H, d, J=8.2Hz), 6.75(2H, d, J=8.9Hz), 4.87(2H, s), 3.85(3H, s); 0.23(9H, s)

**Step 4:** Production of methyl 3-[4-(4'-chloro-4-methoxybiphenyl-2-ylmethoxy)phenylethynyl]-4-cyclohexylaminobenzoate

[4-(4'-Chloro-4-methoxybiphenyl-2-ylmethoxy)phenylethynyl]-trimethylsilane (5.1 g) obtained in the previous step was dissolved in methanol (50 ml) and chloroform (50 ml), and potassium carbonate (2.5 g) was added. The mixture was stirred for 3 hr at room temperature and concentrated. Water (30 ml) was added and the mixture was extracted with ethyl acetate (50 ml). The organic layer was washed with water (30 ml) and saturated brine (30 ml) and dried over sodium sulfate. After filtration, the solvent was evaporated under reduced pressure to give white crystals (3.8 g). The white crystals (2.3 g) were dissolved in acetonitrile (10 ml), and methyl 3-bromo-4-cyclohexylamino-benzoate (1.0 g) obtained in Step 1, tetrakis(triphenylphosphine)palladium complex (0.4 g), copper(I) iodide (0.1 g) and triethylamine (10 ml) were added. The mixture was stirred overnight at 100°C and concentrated under reduced pressure. Water (30 ml) was added and the mixture was extracted with ethyl acetate (50 ml). The organic layer was washed with water (30 ml) and saturated brine (30 ml), and dried over sodium sulfate. After filtration, the solvent was evaporated under reduced pressure and the residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 8:1) to give the title compound (0.9 g, yield 49%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.03(1H, s), 7.84(1H, d, J=8.7Hz), 7.42-7.22(7H, m), 7.15(1H, s), 6.95(1H, d, J=8.2Hz), 6.85(2H, d, J=8.8Hz), 6.59(1H, d, J=8.8Hz), 5.07(1H, brs), 4.91(2H, s), 3.86(3H, s), 3.85(3H, s), 3.42(1H, m), 2.15-2.00(2H, m), 1.80-1.20(8H, m)

**Step 5:** Production of methyl 2-[4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl]-1-cyclohexyl-1H-indole-5-carboxylate

Methyl 3-[4-(4'-chloro-4-methoxybiphenyl-2-ylmethoxy)phenylethynyl]-4-cyclohexylaminobenzoate (0.5 g) obtained in the previous step was dissolved in N,N-dimethylformamide (5 ml), and copper(I) iodide (0.17 g) was added. The mixture was refluxed for

3 hr at 180°C. The insoluble materials were removed by filtration. Water (10 ml) was added and the mixture was extracted with ethyl acetate (30 ml). The organic layer was washed with water (10 ml) and saturated brine (10 ml), and dried over sodium sulfate. After  
5 filtration, the solvent was evaporated under reduced pressure and the residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 8:1) to give the title compound (0.27 g, yield 55%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.34(1H, s), 7.85(1H, d, J=8.8Hz), 7.62(1H, d, J=8.8Hz), 7.40-7.18(8H, m), 7.00-6.94(3H, m), 6.48(1H, s),  
10 4.95(2H, m), 4.18(1H, m), 3.93(3H, s), 3.88(3H, s), 2.45-2.25(2H, m), 1.95-1.20(8H, m)

#### Example 502

Production of 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-  
15 1-cyclohexyl-1H-indole-5-carboxylic acid

Methyl 2-{4-[2-(4-chlorophenyl)-5-methoxybenzyloxy]phenyl}-1-cyclohexyl-1H-indole-5-carboxylate (0.27 g) obtained in Example 501 was treated in the same manner as in Example 2 to give the title compound (0.19 g, yield 71%).

20 APCI-MS: 566 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 12.43(1H, brs), 8.20(1H, s), 7.79(1H, d, J=9.3Hz), 7.72(1H, d, J=9.0Hz), 7.50-7.20(8H, m), 7.07-7.03(3H, m), 6.53(1H, s), 5.01(2H, s), 4.13(1H, m), 3.83(3H, m), 2.35-2.25(2H, m), 1.85-1.10(8H, m)

25 In the same manner as in Examples 501 and 502, and optionally using other conventional methods where necessary, the compound of Example 503 was obtained. The chemical structure and properties are shown in Table 207.

#### Example 601

30 Production of ethyl 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo-[1,2-a]pyridine-7-carboxylate

**Step 1:** Production of 4-benzyloxy-N-methoxy-N-methylbenzamide

4-Benzyloxybenzoic acid (5.0 g) and N,O-dimethylhydroxylamine hydrochloride (2.5 g) were suspended in  
35 dimethylformamide (50 ml), and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (5.0 g), 1-hydroxybenzotriazole (3.5 g) and triethylamine (3.6 ml) were added. The mixture was stirred overnight at room temperature. Water was added to the



reaction mixture and the mixture was extracted with ethyl acetate. The organic layer was washed successively with water, saturated aqueous sodium hydrogencarbonate, water and saturated brine, and dried over anhydrous magnesium sulfate. The solvent was

5 evaporated under reduced pressure to give the title compound (5.6 g, yield 94%).

$^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 7.22, 2H, d,  $J=8.8\text{Hz}$ ), 7.28-7.46 (5H, m), 6.97 (2H, d,  $J=8.8\text{Hz}$ ), 5.10 (2H, s), 3.56 (3H, s), 3.35 (3H, s)

**Step 2:** Production of 1-(4-benzyloxyphenyl)-2-cyclohexylethanone

10 Magnesium (470 mg) was suspended in tetrahydrofuran (2 ml) and cyclohexylmethyl bromide (3.4 g) was added dropwise at room temperature. After the addition, the reaction mixture was stirred for 30 min at  $60^\circ\text{C}$ . The reaction mixture was allowed to cool and diluted with tetrahydrofuran (5 ml). Separately, 4-benzyloxy-N-  
15 methoxy-N-methylbenzamide (3.4 g) obtained in the previous step was dissolved in tetrahydrofuran (10 ml) and the solution was added dropwise to the reaction mixture at room temperature. The mixture was stirred for 2 hr and saturated aqueous ammonium chloride solution was added to the reaction mixture. The mixture  
20 was extracted with diethyl ether. The organic layer was washed with saturated brine and dried over anhydrous magnesium sulfate, and the solvent was evaporated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 9:1) to give the  
25 title compound (3.8 g, yield 66%).

$^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 7.93 (2H, d,  $J=8.8\text{Hz}$ ), 7.28-7.46 (5H, m), 7.00 (2H, d,  $J=8.8\text{Hz}$ ), 5.13 (2H, s), 2.76 (2H, d,  $J=6.8\text{Hz}$ ), 1.95 (1H, m), 0.78-1.82 (10H, m)

**Step 3:** Production of 1-(4-benzyloxyphenyl)-2-bromo-2-

30 cyclohexylethanone

1-(4-Benzyloxyphenyl)-2-cyclohexylethanone (1.0 g) obtained in the previous step was dissolved in 1,4-dioxane (10 ml) and bromine (0.17 ml) was added. The mixture was stirred for 10 min at room temperature. Saturated aqueous sodium hydrogencarbonate  
35 was added to the reaction mixture and the mixture was extracted with diethyl ether. The organic layer was washed with water and saturated brine and dried over anhydrous magnesium sulfate, and the solvent was evaporated under reduced pressure. The residue

was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 9:1) to give the title compound (696 mg, yield 55%).

<sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 7.98(2H, d, J=8.9Hz), 7.28-7.48(5H, m),  
5 7.02(2H, d, J=8.9Hz), 5.14(2H, s), 4.89(1H, d, J=9.3Hz), 0.86-3.30(11H, m)

**Step 4:** Production of ethyl 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo[1,2-a]pyridine-7-carboxylate

Ethyl 2-aminopyridine-4-carboxylate (214 mg) prepared  
10 according to JP-A-8-48651, 1-(4-benzyloxyphenyl)-2-bromo-2-cyclohexylethanone (500 mg) obtained in the previous step and potassium carbonate (356 mg) were stirred for 5 hr with heating at 140°C. The reaction mixture was allowed to cool and chloroform was added. The insoluble materials were filtered off and the  
15 filtrate was concentrated under reduced pressure. The residue was purified by silica gel flash chromatography (developing solvent, n-hexane:ethyl acetate = 1:1) to give the title compound (95 mg, yield 16%).

APCI-MS: 455 (MH<sup>+</sup>)

20 <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 8.33(1H, s), 8.21(1H, d, J=7.5Hz), 7.55(2H, d, J=8.7Hz), 7.25-7.50(6H, m), 5.13(2H, s), 4.41(2H, q, J=7.1Hz), 3.25(1H, m), 1.41(3H, t, J=7.1Hz), 1.15-2.00(10H, m)

#### Example 602

Production of 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo[1,2-a]pyridine-7-carboxylic acid  
25

Ethyl 2-(4-benzyloxyphenyl)-3-cyclohexylimidazo[1,2-a]pyridine-7-carboxylate (95 mg) obtained in the previous step was treated in the same manner as in Example 2 to give the title compound (33 mg, 37%).

30 APCI-MS: 427 (MH<sup>+</sup>)

<sup>1</sup>H-NMR (300MHz, DMSO-d<sub>6</sub>): 8.67(1H, d, J=7.3Hz), 8.08(1H, s), 7.25-7.58(8H, m), 7.13(2H, d, J=8.7Hz), 5.17(2H, s), 3.23(1H, m), 1.25-2.10(10H, m)

The compounds shown in Tables 213 to 218 can be further  
35 obtained in the same manner as in Examples 1 to 701 or by other conventional method employed as necessary.

The evaluation of the HCV polymerase inhibitory activity of the compound of the present invention is explained in the

following. This polymerase is an enzyme coded for by the non-structural protein region called NS5B on the RNA gene of HCV (EMBO J., 15:12-22, 1996).

#### **Experimental Example [I]**

##### **5 i) Preparation of enzyme (HCV polymerase)**

Using, as a template, a cDNA clone corresponding to the full length RNA gene of HCV BK strain obtained from the blood of a patient with hepatitis C, a region encoding NS5B (591 amino acids; J Virol 1991 Mar, 65(3), 1105-13) was amplified by PCR.

10 The objective gene was prepared by adding a 6 His tag {base pair encoding 6 continuous histidine (His)} to the 5' end thereof and transformed to *Escherichia coli*. The *Escherichia coli* capable of producing the objective protein was cultured. The obtained cells were suspended in a buffer solution containing a surfactant and  
15 crushed in a microfluidizer. The supernatant was obtained by centrifugation and applied to various column chromatographys {poly[U]-Sepharese, Sephacryl S-200, mono-S (Pharmacia)}, inclusive of metal chelate chromatography, to give a standard enzyme product.

##### **20 ii) Synthesis of substrate RNA**

Using a synthetic primer designed based on the sequence of HCV genomic 3' untranslated region, a DNA fragment (148 bp) containing polyU and 3'X sequence was entirely synthesized and cloned into plasmid pBluescript SK II(+) (Stratagene). The cDNA  
25 encoding full length NS5B, which was prepared in i) above, was digested with restriction enzyme KpnI to give a cDNA fragment containing the nucleotide sequence of from the restriction enzyme cleavage site to the termination codon. This cDNA fragment was inserted into the upstream of 3' untranslated region of the DNA  
30 in pBluescript SK II(+) and ligated. The about 450 bp inserted DNA sequence was used as a template in the preparation of substrate RNA. This plasmid was cleaved immediately after the 3'X sequence, linearized and purified by phenol-chloroform treatment and ethanol precipitation to give DNA.

35 RNA was synthesized (37°C, 3 hr) by run-off method using this purified DNA as a template, a promoter of pBluescript SK II(+), MEGAscript RNA synthesis kit (Ambion) and T7 RNA polymerase. DNaseI was added and the mixture was incubated for 1

hr. The template DNA was removed by decomposition to give a crude RNA product. This product was treated with phenol-chloroform and purified by ethanol precipitation to give the objective substrate RNA.

5 This RNA was applied to formaldehyde denaturation agarose gel electrophoresis to confirm the quality thereof and preserved at -80°C.

iii) Assay of enzyme (HCV polymerase) inhibitory activity

A test substance (compound of the present invention) and a  
10 reaction mixture (30 µl) having the following composition were reacted at 25°C for 90 min.

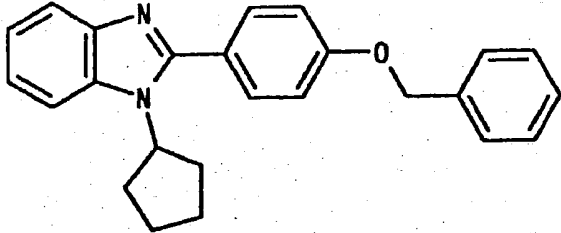
10% Trichloroacetic acid at 4°C and 1% sodium pyrophosphate solution (150 µl) were added to this reaction mixture to stop the reaction. The reaction mixture was left standing in ice for 15  
15 min to insolubilize RNA. This RNA was trapped on a glass filter (Whatman GF/C and the like) upon filtration by suction. This filter was washed with a solution containing 1% trichloroacetic acid and 0.1% sodium pyrophosphate, washed with 90% ethanol and dried. A liquid scintillation cocktail (Packard) was added and  
20 the radioactivity of RNA synthesized by the enzyme reaction was measured on a liquid scintillation counter.

The HCV polymerase inhibitory activity (IC<sub>50</sub>) of the compound of the present invention was calculated from the values of radioactivity of the enzyme reaction with and without the test  
25 substance.

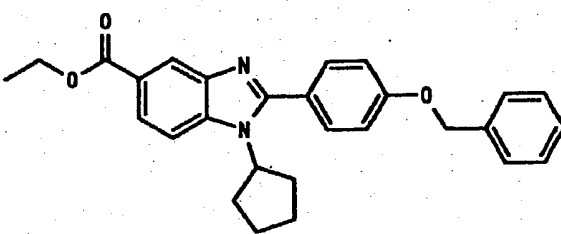
The results are shown in Tables 178-184 and 222-224.

Reaction mixture : HCV polymerase (5 µg/ml) obtained in i), substrate RNA (10 µg/ml) obtained in ii), ATP (50 µM), GTP (50  
30 µM), CTP (50 µM), UTP (2 µM), [5,6-<sup>3</sup>H]UTP (46 Ci/mmol (Amersham), 1.5 µCi) 20 mM Tris-HCl (pH 7.5), EDTA (1 mM), MgCl<sub>2</sub> (5 mM), NaCl (50 mM), DTT (1 mM), BSA (0.01%)

Table 1

Example No.	31	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 7.81 (2H, d, J=6.6Hz), 7.60 (2H, d, J=8.8Hz), 7.51-7.21 (8H, m), 7.11 (2H, d, J=8.8Hz), 5.15 (2H, s), 4.93 (1H, quint, J=8.8Hz), 2.36-2.32 (2H, m), 2.09-2.04 (3H, m), 1.75-1.68 (3H, m).
Purity	> 90% (NMR)	
MS	369 (M+1)	

Example No.	32	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 8.51 (1H, d, J=1.5Hz), 7.98 (1H, d, J=8.4Hz), 7.61 (2H, d, J=8.7Hz), 7.56-7.10 (6H, m), 7.12 (2H, d, J=8.7Hz), 5.15 (2H, s), 4.94 (1H, quint, J=9.3Hz), 4.41 (2H, q, J=7.5Hz), 2.40-1.50 (8H, m), 1.41 (3H, t, J=7.5Hz)
Purity	> 90% (NMR)	
MS	441 (M+1)	

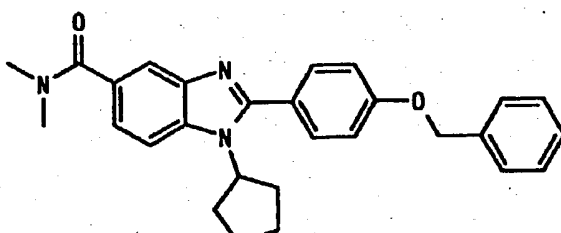
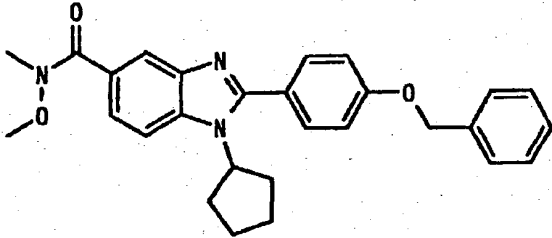
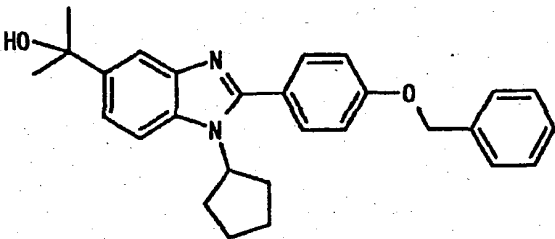
Example No.	33	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 7.84 (1H, s), 7.61 (2H, d, J=9.0Hz), 7.58-7.30 (7H, m), 7.12 (2H, d, J=9.0Hz), 5.15 (2H, s), 4.94 (1H, quint, J=8.7Hz), 3.10 (6H, brs), 2.40-1.50 (8H, m)
Purity	> 90% (NMR)	
MS	440 (M+1)	

Table 2

Example No.	34	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 8.20 (1H, s), 7.50-7.31 (9H, m), 7.12 (2H, d, J=8.7Hz), 5.15 (2H, s), 4.94 (1H, quint, J=8.7Hz), 3.61 (3H, s), 3.40 (3H, s), 2.41-1.42 (8H, m)
Purity	> 90% (NMR)	
MS	456 (M+1)	

Example No.	35	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 7.91 (1H, s), 7.59 (2H, d, J=8.7Hz), 7.49-7.30 (7H, m), 7.11 (2H, d, J=8.8Hz), 5.15 (2H, s), 4.19 (1H, quint, J=8.8Hz), 2.41-2.22 (2H, m), 2.13-1.49 (14H, m)
Purity	> 90% (NMR)	
MS	427 (M+1)	

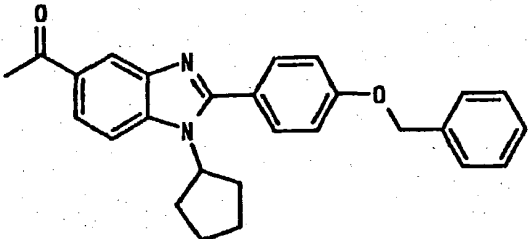
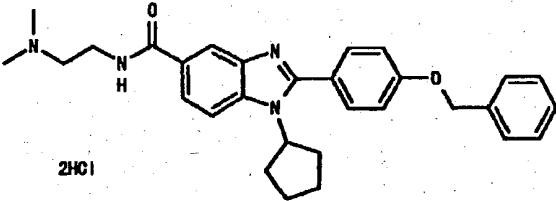
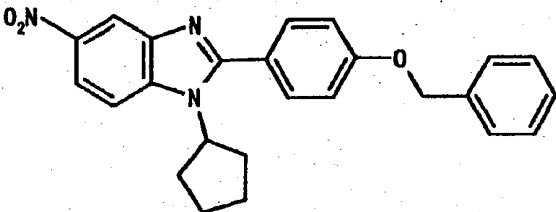
Example No.	36	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 8.40 (1H, d, J=1.4Hz), 7.95 (1H, dd, J=8.6, 1.4Hz), 7.61 (2H, d, J=8.7Hz), 7.57-7.30 (6H, m), 7.13 (2H, d, J=8.7Hz), 5.16 (2H, s), 4.95 (1H, quint, J=8.8Hz), 2.64 (3H, s), 2.40-1.54 (8H, m)
Purity	> 90% (NMR)	
MS	411 (M+1)	

Table 3

Example No.	37	1H NMR ( $\delta$ ) ppm
 <p>2HCl</p>		300MHz, DMSO-d <sub>6</sub> 10.47 (1H, brs, ), 9.15 (1H, brs), 8.40 (1H, s), 8.07 (1H, d, J=9.0Hz), 7.93 (1H, d, J=8.7Hz), 7.77 (2H, d, J=8.7Hz), 7.55-7.29 (7H, m), 5.26 (2H, s), 4.93 (1H, quint, J=9.0Hz), 3.77-3.63 (2H, m), 3.39-3.23 (2H, m), 2.84 (6H, d, J=4.8Hz), 2.32-1.60 (8H, m)
Purity	> 90% (NMR)	
MS	483 (M+1)	

Example No.	38	1H NMR ( $\delta$ ) ppm
		300MHz, CDCl <sub>3</sub> 8.69 (1H, s), 8.19 (1H, d, J=9.0Hz), 7.62 (2H, d, J=8.7Hz), 7.54 (1H, d, J=9.0Hz), 7.48-7.36 (5H, m), 7.15 (2H, d, J=8.7Hz), 5.17 (2H, s), 4.98 (1H, quint, J=9.0Hz), 2.27-2.07 (6H, m), 1.82-1.78 (2H, m)
Purity	> 90% (NMR)	
MS	414 (M+1)	

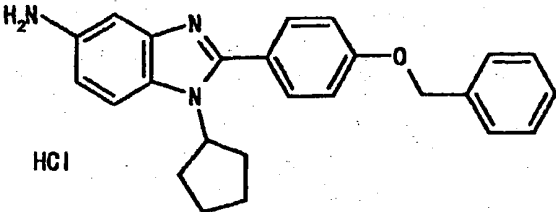
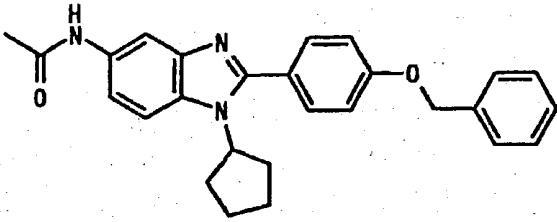
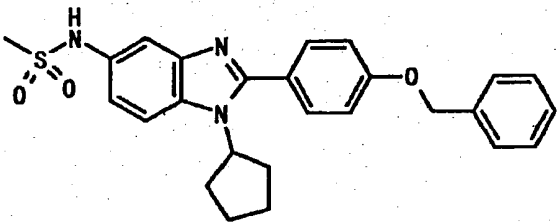
Example No.	39	1H NMR ( $\delta$ ) ppm
 <p>HCl</p>		300MHz, DMSO-d <sub>6</sub> 7.84 (1H, d, J=9.0Hz), 7.79 (2H, d, J=8.7Hz), 7.52-7.33 (8H, m), 7.26 (1H, d, J=9.0Hz), 5.27 (2H, s), 4.92 (1H, quint, J=9.3Hz), 2.19-1.70 (8H, m)
Purity	> 90% (NMR)	
MS	384 (M+1)	

Table 4

Example No.	40	1H NMR(δ) ppm
		300MHz, CDCl <sub>3</sub> 7.72(1H, s), 7.60-7.35(10H, m), 7.10(2H, d, J=8.7Hz), 5.14(2H, s), 4.90(1H, quint, J=8.8Hz), 2.29-2.19(2H, m), 2.19(3H, s), 2.19-1.74(6H, m).
Purity	> 90% (NMR)	
MS	426(M+1)	

Example No.	41	1H NMR(δ) ppm
		300MHz, CDCl <sub>3</sub> 7.66(1H, s), 7.61(2H, d, J=8.8Hz), 7.50-7.28(7H, m), 7.12(2H, d, J=8.8Hz), 6.86(1H, brs), 5.15(2H, s), 4.94(1H, quint, J=8.8Hz), 2.97(3H, s), 2.29-1.76(8H, m).
Purity	> 90% (NMR)	
MS	462(M+1)	

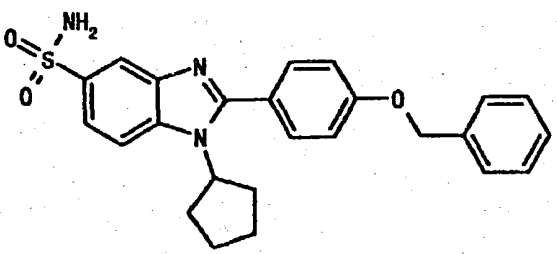
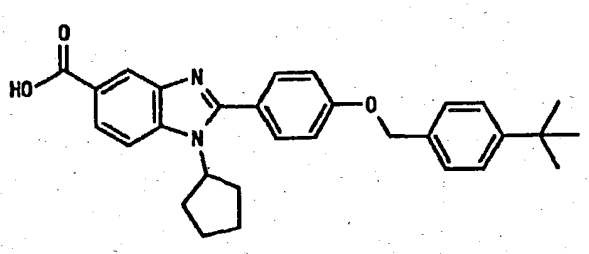
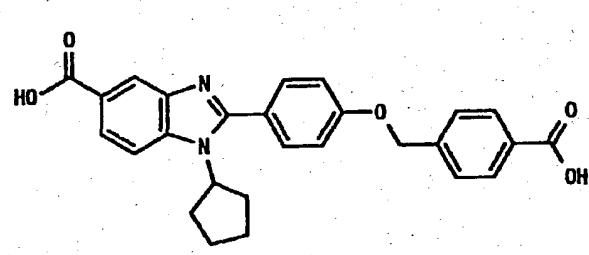
Example No.	42	1H NMR(δ) ppm
		300MHz, DMSO 8.11(1H, s), 7.81(1H, d, J=8.4Hz), 7.72(1H, d, J=8.4Hz), 7.65(2H, d, J=8.4Hz), 7.51(2H, m), 7.43(2H, m), 7.37(1H, m), 7.29(2H, s), 7.23(2H, d, J=8.4Hz), 5.22(2H, s), 4.89(1H, quintet, J=9.2Hz), 2.2-2.0(6H, m), 1.7(2H, m).
Purity	> 90% (NMR)	
MS	448(M+)	



Table 5

Example No.	43	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.08 (1H, d, J=9.0Hz), 7.99 (1H, d, J=9.0Hz), 7.47-7.41 (4H, m), 7.33 (2H, d, J=8.4Hz), 5.22 (2H, s), 4.96 (1H, quint, J=9.0Hz), 2.25-1.60 (8H, m), 1.30 (9H, s)
Purity	> 90% (NMR)	
MS	469 (M+1)	

Example No.	44	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.9 (2H, brs), 8.25 (1H, s), 8.00 (2H, d, J=7.8Hz), 7.90 (1H, d, J=8.4Hz), 7.74 (1H, d, J=8.7Hz), 7.67 (2H, d, J=9.0Hz), 7.62 (2H, d, J=8.1Hz), 7.24 (2H, d, J=8.4Hz), 5.32 (2H, s), 4.88 (1H, quint, J=9.0Hz), 2.25-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	457 (M+1)	

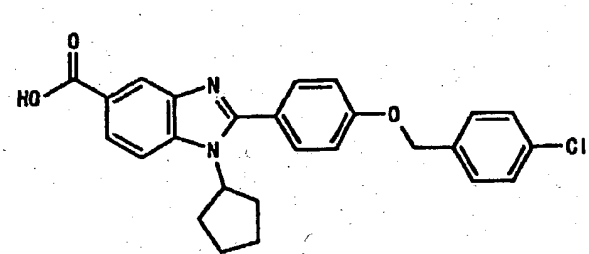
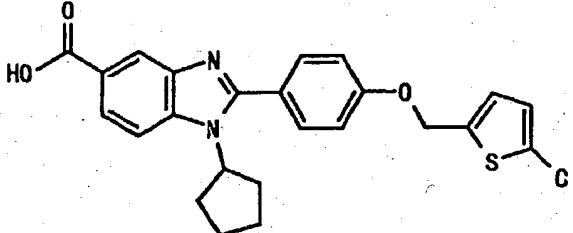
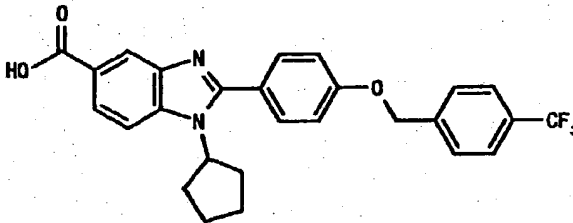
Example No.	45	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.4 (1H, brs), 8.32 (1H, s), 8.06 (1H, d, J=8.7Hz), 7.97 (1H, d, J=8.7Hz), 7.79 (2H, d, J=8.8Hz), 7.56-7.48 (4H, m), 7.33 (2H, d, J=8.8Hz), 5.27 (2H, s), 4.95 (1H, quint, J=8.9Hz), 2.30-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	447 (M+1)	

Table 6

Example No.	46	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.07 (1H, d, J=8.7Hz), 7.98 (1H, d, J=8.7Hz), 7.80 (2H, d, J=8.4Hz), 7.34 (2H, d, J=8.4Hz), 7.19 (1H, d, J=3.6Hz), 7.09 (1H, d, J=3.6Hz), 5.41 (2H, s), 4.95 (1H, quint, J=8.7Hz), 2.30-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	453 (M+1)	

Example No.	47	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.07 (1H, d, J=8.4Hz), 7.98 (1H, d, J=9.0Hz), 7.82-7.72 (6H, m), 7.35 (2H, d, J=9.0Hz), 5.40 (2H, s), 4.95 (1H, quint, J=8.7Hz), 2.35-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	481 (M+1)	

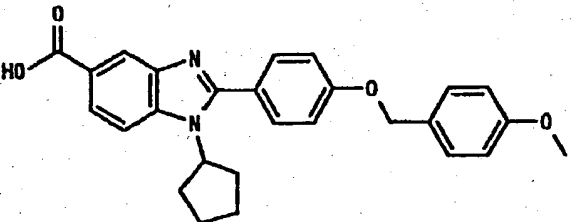
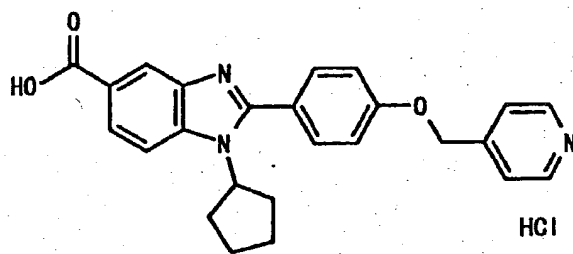
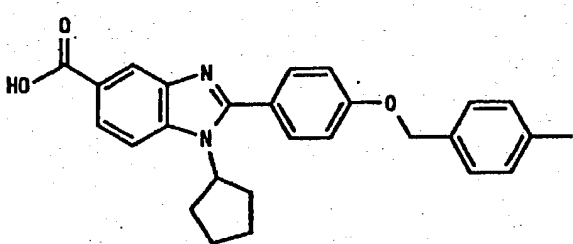
Example No.	48	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.88 (1H, d, J=8.4Hz), 7.70 (1H, d, J=8.4Hz), 7.64 (2H, d, J=8.4Hz), 7.43 (2H, d, J=8.4Hz), 7.20 (2H, d, J=8.4Hz), 6.98 (2H, d, J=8.4Hz), 5.13 (2H, s), 4.88 (1H, quint, J=8.7Hz), 3.77 (3H, s), 2.35-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	443 (M+1)	

Table 7

Example No.	49	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.93 (2H, d, J=6.6Hz), 8.35 (1H, s), 8.06-8.04 (3H, m), 7.97 (1H, d, J=8.7Hz), 7.83 (2H, d, J=8.7Hz), 7.38 (2H, d, J=8.7Hz), 5.61 (2H, s), 4.94 (1H, quint, J=8.7Hz), 2.40-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	414 (M+1)	

Example No.	50	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.08 (1H, d, J=8.7Hz), 7.99 (1H, d, J=9.0Hz), 7.78 (2H, d, J=8.4Hz), 7.39 (2H, d, J=8.1Hz), 7.32 (2H, d, J=8.7Hz), 7.23 (2H, d, J=7.8Hz), 5.22 (2H, s), 4.96 (1H, quint, J=9.0Hz), 2.32 (3H, s), 2.30-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	427 (M+1)	

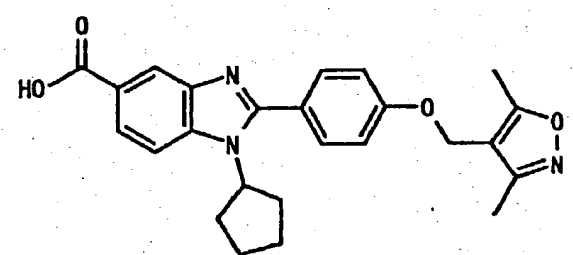
Example No.	51	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.03 (1H, d, J=9.0Hz), 7.93 (1H, d, J=9.0Hz), 7.77 (2H, d, J=8.4Hz), 7.31 (2H, d, J=8.7Hz), 5.07 (2H, s), 4.94 (1H, quint, J=8.7Hz), 2.45 (3H, s), 2.26 (3H, s), 2.26-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	432 (M+1)	

Table 8

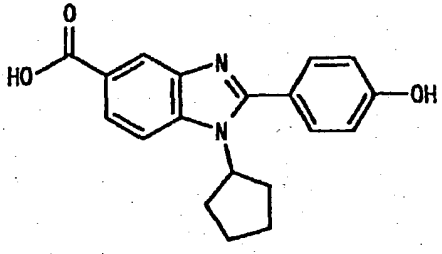
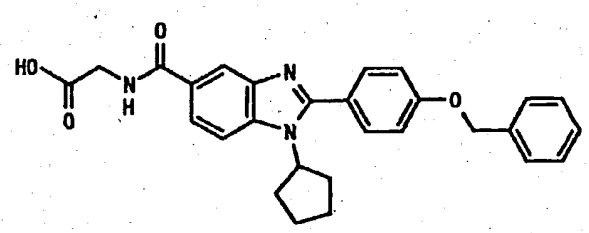
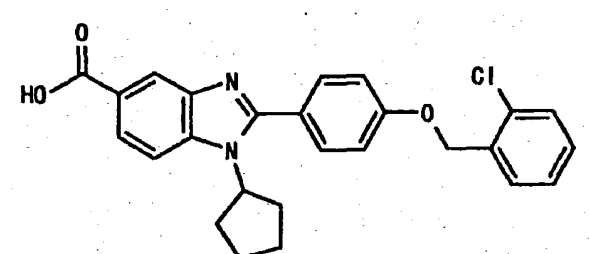
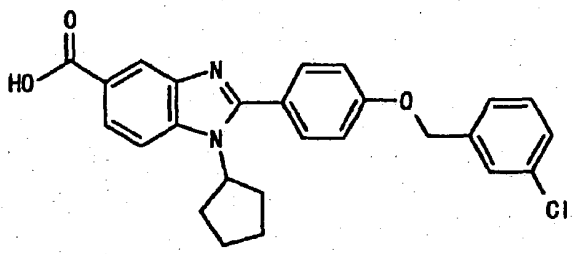
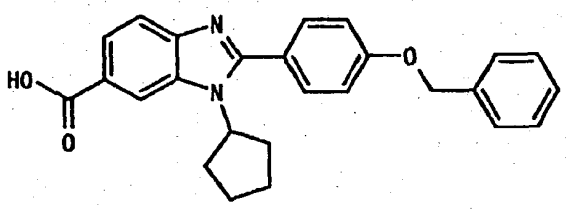
Example No.	52	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.7 (1H, brs), 10.0 (1H, s), 8.22 (1H, s), 7.87 (1H, d, J=8.6Hz), 7.69 (1H, d, J=8.6Hz), 7.53 (2H, d, J=8.6Hz), 6.96 (2H, d, J=8.6Hz), 4.89 (1H, q, J=9.0Hz), 2.30-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	323 (M+1)	
Example No.	53	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9.18 (1H, t, J=5.6Hz), 8.34 (1H, s), 8.04 (1H, d, J=9.6Hz), 7.98 (1H, d, J=8.7Hz), 7.80 (2H, d, J=8.7Hz), 7.52-7.32 (7H, m), 5.27 (2H, s), 4.95 (1H, quint, J=9.0Hz), 3.99 (2H, d, J=5.7Hz), 2.40-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	470 (M+1)	
Example No.	54	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.32 (1H, s), 8.05 (1H, d, J=8.7Hz), 7.95 (1H, d, J=8.7Hz), 7.80 (2H, d, J=8.4Hz), 7.67 (1H, t, J=4.5Hz), 7.56 (1H, t, J=4.5Hz), 7.45-7.42 (2H, m), 7.35 (2H, d, J=8.4Hz), 5.31 (2H, s), 4.96 (1H, quint, J=9.0Hz), 2.30-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	447 (M+1)	

Table 9

Example No.	55	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.78 (1H, br s), 8.24 (1H, s), 7.88 and 7.72 (2H, ABq, J=8.6Hz), 7.66 and 7.23 (4H, A'B'q, J=8.6Hz), 7.58 (1H, s), 7.48-7.42 (3H, m), 5.24 (1H, s), 4.88 (1H, quint, J=8.8Hz), 2.30-1.91 (6H, m), 1.78-1.60 (2H, m)
Purity	> 90% (NMR)	
MS	447 (M+1)	

Example No.	56	1H NMR (δ) ppm
		300MHz, DMSO 12.89 (1H, broad), 8.18 (1H, s), 7.87 (1H, d, J=8.4Hz), 7.74 (1H, d, J=9.2Hz), 7.67 (2H, d, J=8.8Hz), 7.52 (2H, m), 7.45 (2H, m), 7.38 (1H, m), 7.23 (2H, d, J=8.8Hz), 5.22 (2H, s), 4.94 (1H, quintet, J=8.9Hz), 2.16 (4H, m), 1.98 (2H, m), 1.73 (2H, m).
Purity	> 90% (NMR)	
MS	413 (M+)	

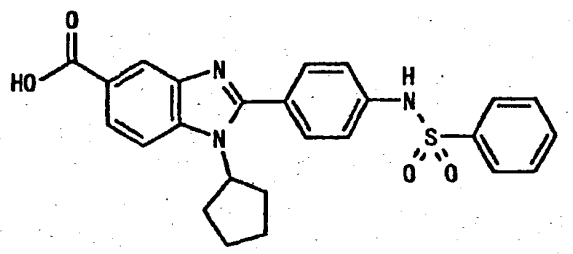
Example No.	57	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.99 (1H, s), 8.26 (1H, s), 8.01-7.86 (4H, m), 7.69-7.59 (5H, m), 7.38 (2H, d, J=8.7Hz), 4.86 (1H, quint, J=8.7Hz), 2.12-1.90 (6H, m), 1.72-1.59 (2H, m)
Purity	> 90% (NMR)	
MS	462 (M+1)	

Table 10

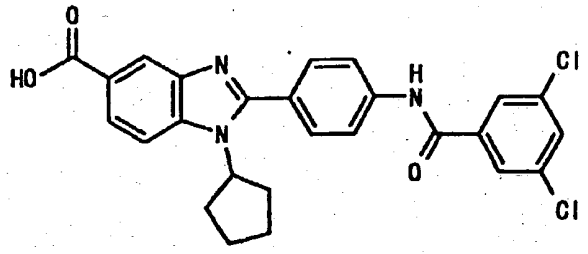
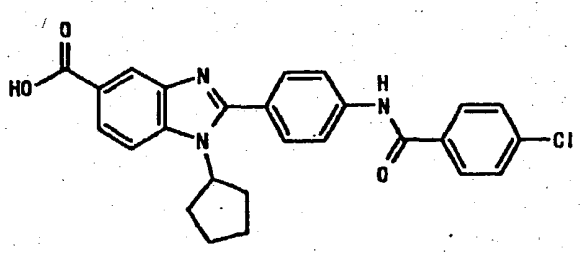
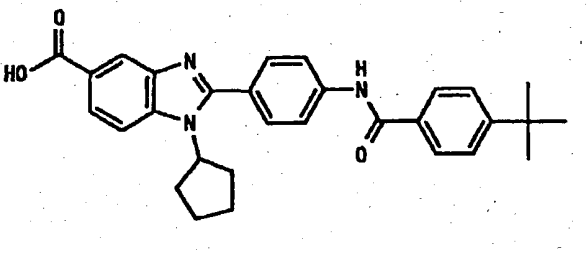
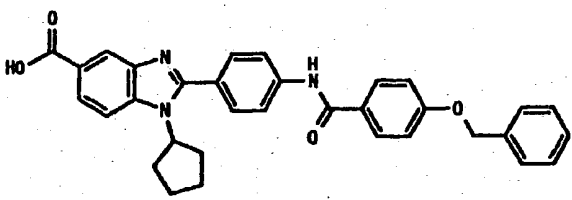
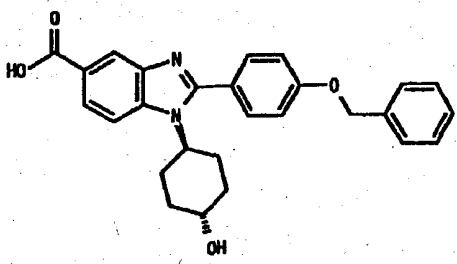
Example No.	58	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.78 (1H, s), 10.69 (1H, s), 8.26-7.72 (9H, m), 4.92 (1H, quint, J=9.0Hz), 2.34-1.70 (6H, m), 1.75-1.61 (2H, m)
Purity	> 90% (NMR)	
MS	494 (M+1)	
Example No.	59	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.82 (1H, s), 8.34 (1H, s), 8 .14 and 7.84 (4H, ABq, J=8.4H z), 8.06 and 7.66 (4H, A' B' q, J=8.6Hz), 8.06-7.98 (4H, m) , 5.01 (1H, quint, J=9.3Hz), 2.35-2.15 (4H, m), 2.11-1.9 6 (2H, m), 1.80-1.62 (2H, m)
Purity	> 90% (NMR)	
MS	460 (M+1)	
Example No.	60	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.61 (1H, s), 8.32 (1H, s), 8 .12 and 7.81 (4H, ABq, J=8.9H z), 8.03 and 7.93 (2H, A' B' q, J=8.7Hz), 7.95 and 7.59 (4H, A'' B'' q, J=8.4Hz), 4.99 (1H, q uint, J=9.0Hz), 2.33-2.12 ( 4H, m), 2.10-1.93 (2H, m), 1. 80-1.63 (2H, m), 1.34 (9H, m)
Purity	> 90% (NMR)	
MS	482 (M+1)	

Table 11

Example No.	61	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.6 (1H, s), 8.34 (1H, s), 8.13 (2H, d, J=8.7Hz), 8.09-7.98 (4H, m), 7.82 (2H, d, J=8.7Hz), 7.50-7.35 (5H, m), 7.20-7.17 (2H, d, J=9.0Hz), 5.24 (2H, s), 5.01 (1H, quint, J=9.3Hz), 2.40-1.60 (8H, m).
Purity	> 90% (NMR)	
MS	532 (M+1)	

Example No.	62	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.26 (1H, d, J=8.7Hz), 8.04 (1H, d, J=8.7Hz), 7.77 (2H, d, J=8.4Hz), 7.52 (2H, d, J=6.9Hz), 7.46-7.39 (5H, m), 5.28 (2H, s), 4.38 (1H, m), 3.71 (1H, m), 2.60-2.15 (2H, m), 2.04-1.96 (4H, m), 1.30-1.20 (2H, m).
Purity	> 90% (NMR)	
MS	443 (m+1)	

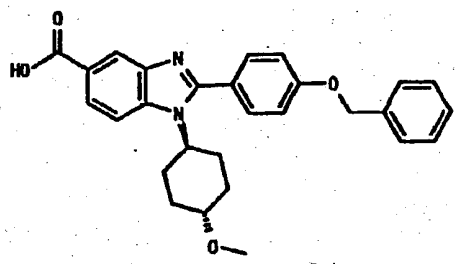
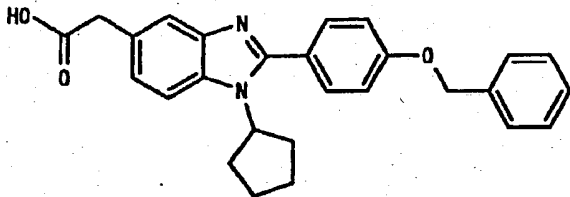
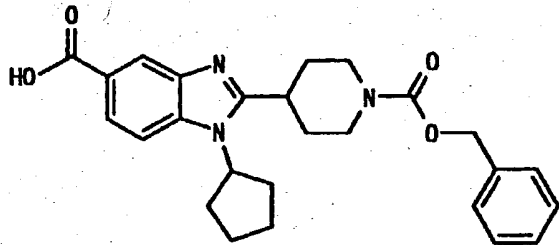
Example No.	63	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 8.14 (1H, d, J=8.7Hz), 7.96 (1H, d, J=8.4Hz), 7.71 (2H, d, J=9.0Hz), 7.51 (2H, d, J=6.9Hz), 7.46-7.37 (3H, m), 7.30 (2H, d, J=8.4Hz), 5.25 (3H, s), 4.39 (1H, m), 3.44 (1H, m), 3.27 (3H, s), 2.60-1.95 (6H, m), 1.25-1.05 (2H, m).
Purity	約 90% (NMR)	
MS	457 (M+1)	

Table 12

Example No.	64	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.25 (1H, brs), 7.70-7.30 (9H, m), 7.20 (2H, d, J=8.7Hz), 7.14 (1H, d, J=8.4Hz), 5.20 (2H, s), 4.84 (1H, quint, J=6.0Hz), 3.66 (2H, s), 2.30-1.51 (8H, m)
Purity	> 90% (NMR)	
MS	427 (M+1)	

Example No.	65	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.64 (1H, brs), 8.13 (1H, s), 7.80 (1H, d, J=7.2Hz), 7.59 (1H, d, J=8.7Hz), 7.48-7.30 (5H, m), 5.11 (2H, s), 5.03 (1H, quint, J=8.7Hz), 4.20-4.05 (2H, m), 3.45-3.90 (3H, m), 2.15-1.60 (12H, m)
Purity	> 90% (NMR)	
MS	448 (M+1)	

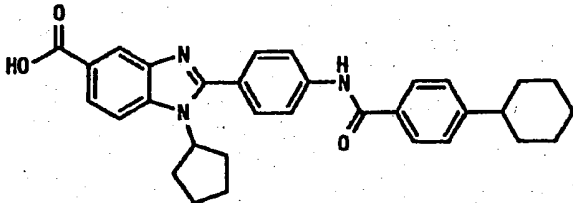
Example No.	66	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.59 (1H, s), 8.31 (1H, s), 8.10 (2H, d, J=8.6Hz), 8.03 (1H, d, J=8.7Hz), 8.00-7.85 (3H, m), 7.80 (2H, d, J=8.6Hz), 7.41 (2H, d, J=8.2Hz), 4.98 (1H, quint, J=8.8Hz), 2.71-1.10 (19H, m)
Purity	> 90% (NMR)	
MS	508 (M+1)	



Table 13

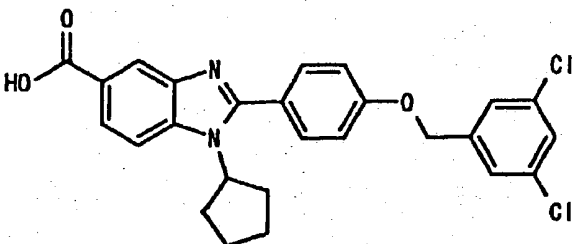
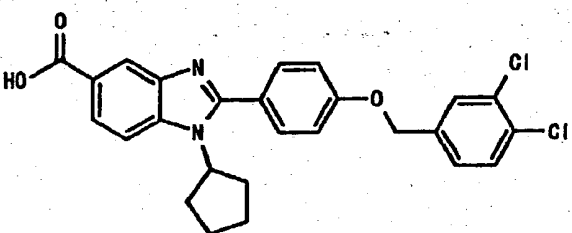
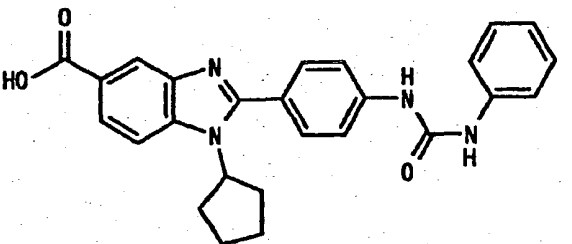
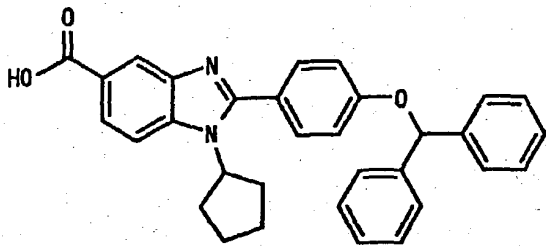
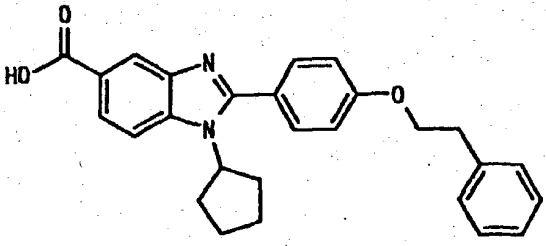
Example No.	67	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.81 (1H, brs), 8.42 (1H, s), 7.90 (1H, d, J=8.5Hz), 7.80 -7.52 (6H, m), 7.44 (2H, d, J= 8.6Hz), 5.25 (2H, s), 4.88 (1 H, quint, J=8.8Hz), 2.30-1. 52 (8H, m)
Purity	> 90% (NMR)	
MS	481 (M+1)	
Example No.	68	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, d, J=1.4Hz), 8.05 (1H, d, J=8.6Hz), 7.96 (1H, d, J=8.6Hz), 8.86-8.61 (4H, m), 7.51 (1H, d, J=6.3Hz), 7.33 (2H, d, J=8.8Hz), 5.28 (2H, s), 4.94 (1H, quint, J=8.8Hz), 2.31-1.60 (8H, m)
Purity	> 90% (NMR)	
MS	481 (M+1)	
Example No.	69	1H NMR (δ) ppm
		300MHz, DMSO-d6 9.88 (1H, s), 9.42 (1H, s), 8.32 (1H, s), 8.09 and 8.02 (2H, ABq, J=9.0Hz), 7.81 and 7.78 (4H, A'B'q, J=9.2Hz), 7.50 (2H, d, J=7.8Hz), 7.31 (2H, t, J=7.8Hz), 7.00 (1H, t, J=7.8Hz), 5.03 (1H, quint, J=8.7Hz), 2.34-2.17 (4H, m), 2.13-1.96 (2H, m), 1.83-1.64 (2H, m)
Purity	> 90% (NMR)	
MS	441 (M+1)	

Table 14

Example No.	70	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, d, J=1.2Hz), 8.04 (1H, d, J=8.7Hz), 7.94 (1H, d, J=8.7Hz), 7.72 (2H, d, J=8.7Hz), 7.60-7.20 (12H, m), 6.74 (1H, s), 4.92 (1H, quint, J=8.9Hz), 2.30-1.58 (8H, m)
Purity	> 90% (NMR)	
MS	489 (M+1)	

Example No.	71	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.05 (1H, d, J=8.7Hz), 7.97 (1H, d, J=8.7Hz), 7.76 (2H, d, J=8.6Hz), 7.44-7.19 (7H, m), 4.94 (1H, quint, J=8.8Hz), 4.35 (2H, t, J=6.7Hz), 3.10 (2H, t, J=6.7Hz), 2.32-1.60 (8H, m)
Purity	> 90% (NMR)	
MS	427 (M+1)	

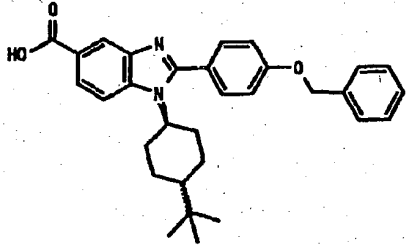
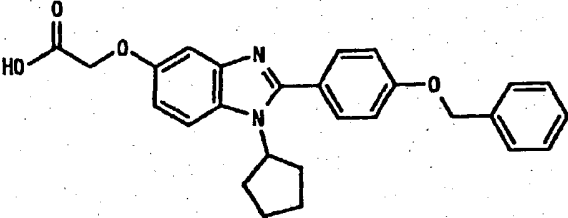
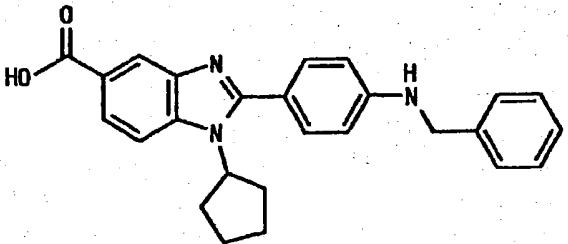
Example No.	72	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.25 (1H, d, J=8.7Hz), 8.03 (1H, d, J=9.0Hz), 7.75 (2H, d, J=8.7Hz), 7.51 (2H, d, J=7.2Hz), 7.46-7.33 (5H, m), 5.27 (2H, s), 4.36 (1H, m), 2.50-2.25 (2H, m), 2.15-2.00 (2H, m), 1.95-1.85 (2H, m), 1.35 (1H, m), 1.20-1.10 (2H, m), 0.87 (9H, s)
Purity	> 90% (NMR)	
MS	483 (M+1)	

Table 15

Example No.	73	1H NMR (δ) ppm
		300MHz, DMSO-d6 7.59 (2H, d, J=8.4Hz), 7.52-7.35 (6H, m), 7.20 (2H, d, J=8.7Hz), 7.14 (1H, d, J=2.1Hz), 6.90 (1H, dd, J=9.0, 2.4Hz), 5.21 (2H, s), 4.83 (1H, quint, J=8.7Hz), 4.70 (2H, s), 2.30-1.90 (6H, m), 1.75-1.55 (2H, m).
Purity	> 90% (NMR)	
MS	443 (M+1)	

Example No.	74	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 8.06 and 7.97 (2H, ABq, J=8.7Hz), 7.57 and 6.86 (4H, A'B'q, J=8.9Hz), 7.42-7.26 (5H, m), 5.04 (1H, quint, J=9.0Hz), 4.42 (2H, s), 2.32-1.94 (6H, m), 1.80-1.62 (2H, m)
Purity	> 90% (NMR)	
MS	412 (M+1)	

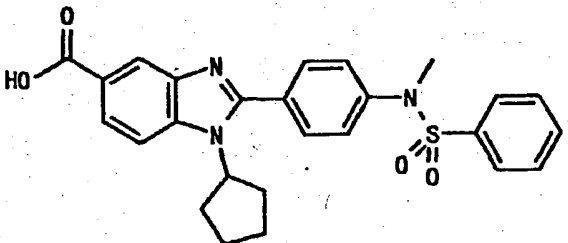
Example No.	75	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.80 (1H, s), 8.26 (1H, s), 7.90 (1H, d, J=9.2Hz), 7.76-7.60 (8H, m), 7.35 (2H, d, J=8.4Hz), 4.84 (1H, quint, J=8.8Hz), 3.23 (3H, s), 2.32-1.90 (6H, m), 1.78-1.61 (2H, m)
Purity	> 90% (NMR)	
MS	476 (M+1)	

Table 16

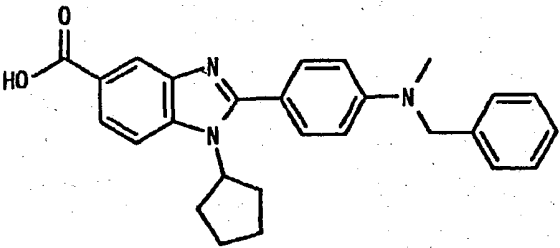
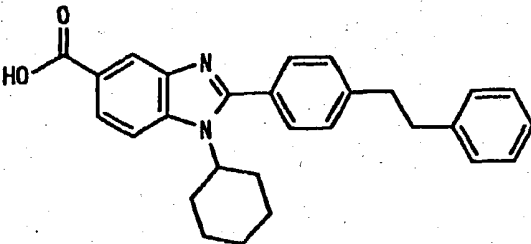
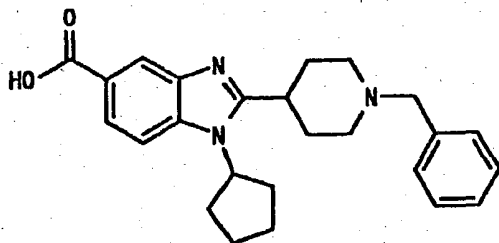
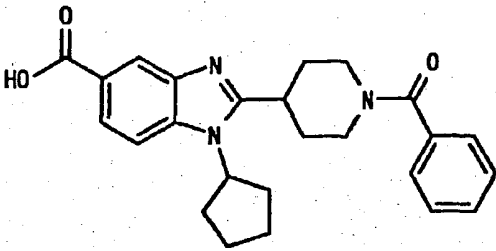
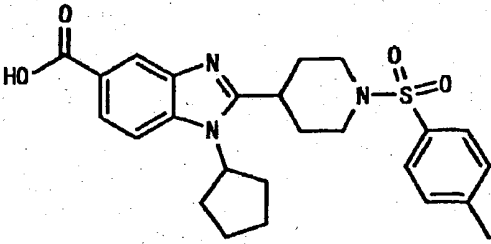
Example No.	76	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.07 and 7.49 (2H, ABq, J=8.7Hz), 7.66 and 7.00 (4H, A'B'q, J=7.7Hz), 7.39-7.24 (5H, m), 5.05 (1H, quint, J=8.8Hz), 4.76 (2H, s), 3.21 (3H, s), 2.35-1.92 (6H, m), 1.81-1.62 (2H, m)
Purity	> 90% (NMR)	
MS	426 (M+1)	
Example No.	77	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 7.87 (1H, s), 7.56 and 7.43 (4H, ABq, J=8.1Hz), 7.34-7.16 (5H, m), 4.25 (1H, brt, J=12.5Hz), 3.06-2.92 (4H, m), 2.41-2.17 (2H, m), 1.96-1.77 (4H, m), 1.72-1.58 (1H, m), 1.48-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	425 (M+1)	
Example No.	78	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.14 (1H, s), 7.79 (1H, d, J=9.0Hz), 7.57 (1H, d, J=8.7Hz), 7.40-7.20 (5H, m), 4.89 (1H, quint, J=8.7Hz), 3.54 (2H, s), 3.19-2.90 (3H, m), 2.23-1.69 (14H, m)
Purity	> 90% (NMR)	
MS	404 (M+1)	

Table 17

Example No.	79	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.15 (1H, s), 7.81 (1H, d, J=8.4Hz), 7.59 (1H, d, J=9.0Hz), 7.50-7.38 (5H, m), 5.05 (1H, quint, J=9.0Hz), 3.85-2.95 (3H, m), 2.20-1.65 (14H, m)
Purity	> 90 % (NMR)	
MS	418 (M+1)	

Example No.	80	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.17 (1H, m), 7.84 (1H, d, J=8.4Hz), 7.78-7.62 (3H, m), 7.49 (2H, d, J=8.1Hz), 5.05-4.91 (1H, m), 3.80-3.70 (2H, m), 3.30-3.12 (1H, m), 2.48-2.31 (5H, m), 2.15-1.60 (12H, m)
Purity	> 90 % (NMR)	
MS	468 (M+1)	

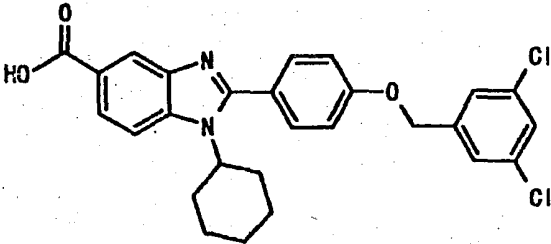
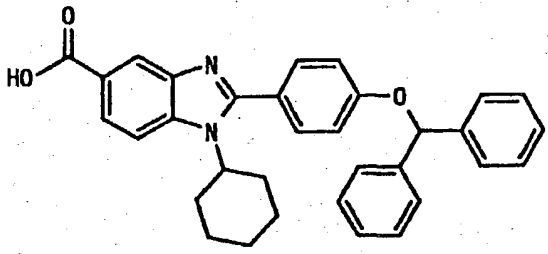
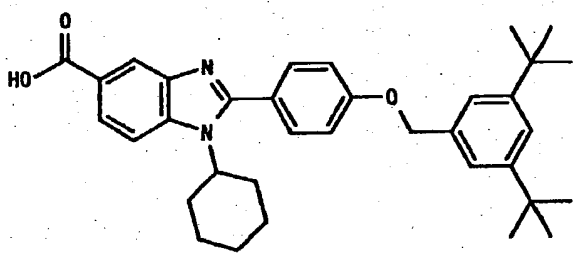
Example No.	81	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, brs), 8.21 (1H, d, J=1.4Hz), 7.49 (1H, d, J=8.6Hz), 7.85 (1H, dd, J=8.6, 1.4Hz), 7.70-7.55 (5H, m), 7.23 (2H, d, J=8.7Hz), 5.25 (2H, s), 4.36-4.15 (1H, m), 2.39-2.18 (2H, m), 2.00-1.78 (4H, m), 1.70-1.57 (1H, m), 1.48-1.15 (3H, m)
Purity	> 90 % (NMR)	
MS	495 (M+1)	

Table 18

Example No.	82	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 8.22 (1H, d, J=8.7Hz), 8.02 (1H, d, J=8.7Hz), 7.69 (2H, d, J=8.7Hz), 7.60-7.50 (4H, m), 7.45-7.25 (8H, m), 6.75 (1H, s), 4.21-4.23 (1H, m), 2.39-2.18 (2H, m), 2.10-1.78 (4H, m), 1.70-1.15 (4H, m)
Purity	> 90% (NMR)	
MS	503 (M+1)	

Example No.	83	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.2 (1H, brs), 8.30 (1H, s), 8.23 (1H, d, J=8.8Hz), 8.02 (1H, d, J=8.7Hz), 7.74 (2H, d, J=8.6Hz), 7.40-7.33 (5H, m), 5.22 (2H, s), 4.36 (1H, m), 2.50-1.40 (10H, m), 1.31 (18H, s).
Purity	> 90% (NMR)	
MS	539 (M+1)	

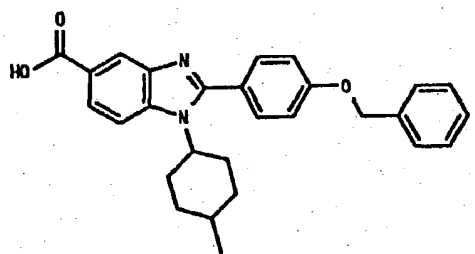
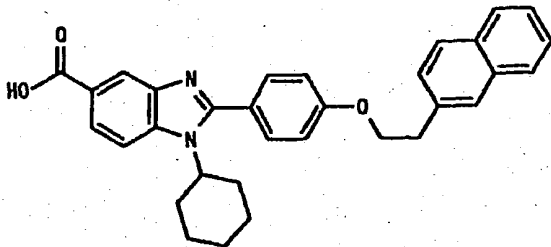
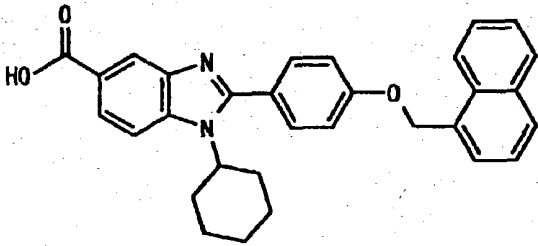
Example No.	84	1H NMR (δ) ppm
		mixture of isomers (cis:trans=3:1) 300MHz, DMSO-d6 8.30 (1H, s), 8.20-7.95 (2H, m), 7.72 (2H, d, J=8.4Hz), 7.52-7.29 (7H, m), 5.25 (2H, s), 4.34, 3.40 (1H, m), 2.50-2.20 (2H, m), 2.05-1.50 (6H, m), 1.14, 0.90 (3H, d, J=6.9, 6.3Hz), 1.09 (1H, m).
Purity	> 90% (NMR)	
MS	441 (M+1)	

Table 19

Example No.	85	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.25 (1H, s), 8.14-7.83 (6H, m), 7.77-7.44 (5H, m), 7.21 (2H, d, J=7.8Hz), 4.44 (2H, brt), 4.31 (1H, brt), 3.56 (2H, brt), 2.20-2.16 (2H, m), 2.00-1.74 (4H, m), 1.70-1.55 (1H, m), 1.45-1.14 (3H, m)
Purity	> 90% (NMR)	
MS	491 (M+1)	

Example No.	86	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, s), 8.23 (1H, s), 8.15 (1H, d, J=7.6Hz), 8.02-7.53 (10H, m), 7.32 (2H, d, J=8.7Hz), 5.68 (2H, s), 4.32 (1H, brt, J=12.2Hz), 2.41-2.20 (2H, m), 2.01-1.78 (4H, m), 1.71-1.56 (1H, m), 1.50-1.16 (3H, m)
Purity	> 90% (NMR)	
MS	477 (M+1)	

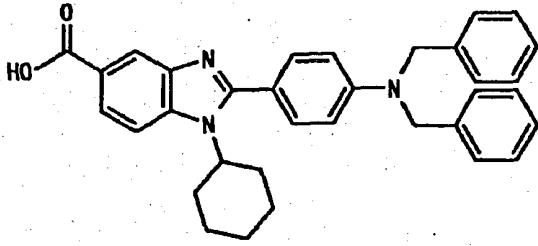
Example No.	87	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, brs), 8.16 (1H, s), 7.91 and 7.82 (2H, ABq, J=8.5Hz), 7.44 and 6.86 (4H, A'B'q, J=8.6Hz), 7.39-7.26 (10H, m), 4.82 (2H, s), 4.35 (1H, brt, J=12.2Hz), 2.35-2.16 (2H, m), 1.97-1.75 (4H, m), 1.69-1.56 (1H, m), 1.45-1.16 (3H, m)
Purity	> 90% (NMR)	
MS	516 (M+1)	

Table 20

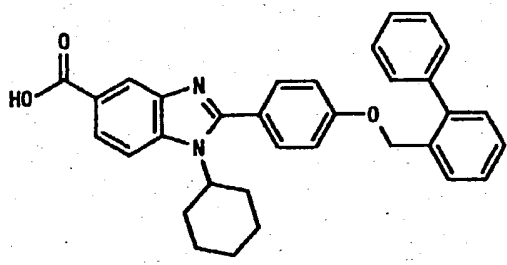
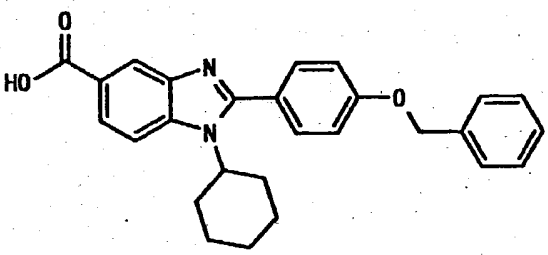
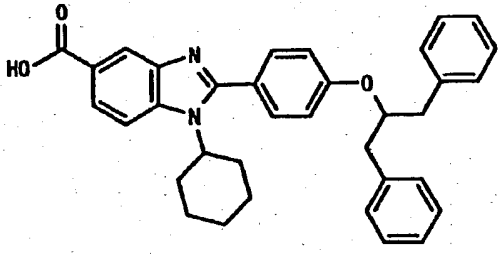
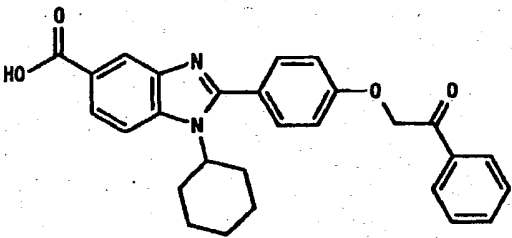
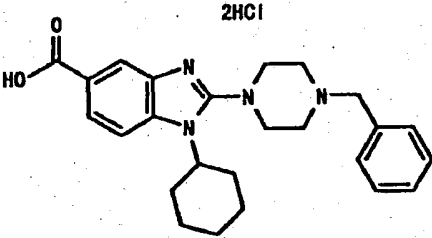
Example No.	88	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.26 and 8.06 (2H, ABq, J=8.9Hz), 7.73 and 7.22 (4H, A'B'q, J=8.7Hz), 7.50-7.36 (8H, m), 5.10 (2H, s), 4.37 (1H, brt, J=12.2Hz), 2.38-2.28 (2H, m), 2.10-1.80 (4H, m), 1.70-1.56 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	503 (M+1)	
Example No.	89	1H NMR (δ) ppm
		
Purity	91% (HPLC)	
MS	427 (M+1)	
Example No.	90	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.40-8.20 (2H, m), 8.04 (1H, d, J=8.4Hz), 7.65 (2H, d, J=8.4Hz), 7.50-7.10 (12H, m), 5.08 (1H, m), 4.33 (1H, m), 3.00 (4H, m), 2.50-1.10 (10H, m)
Purity	> 90% (NMR)	
MS	531 (M+1)	



Table 21

Example No.	91	1H NMR ( $\delta$ ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.27 (1H, d, J=8.7Hz), 8.08-8.03 (3H, m), 7.77-7.58 (5H, m), 7.31 (2H, d, J=8.7Hz), 5.81 (2H, s), 4.40 (1H, m), 2.50-1.20 (10H, m).
Purity	約90% (NMR)	
MS	455 (M+1)	

Example No.	92	1H NMR ( $\delta$ ) ppm
		300MHz, DMSO-d6 11.8 (1H, brs), 8.07 (1H, s), 7.89 (1H, d, J=8.7Hz), 7.84 (1H, d, J=8.4Hz), 7.69 (2H, m), 7.48 (3H, m), 4.42 (2H, s), 4.11 (1H, m), 3.73 (4H, m), 3.40 (4H, m), 2.40-1.40 (10H, m).
Purity	>90% (NMR)	
MS	419 (M+1)	

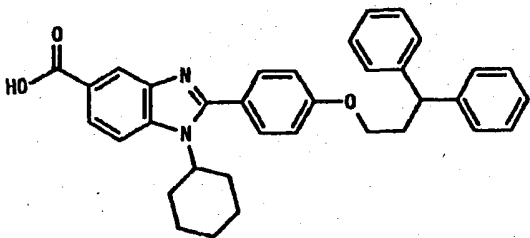
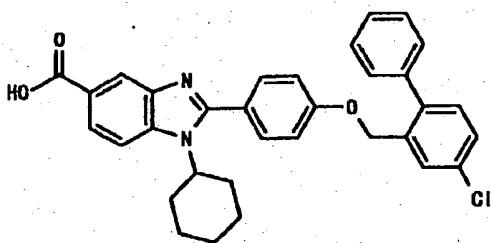
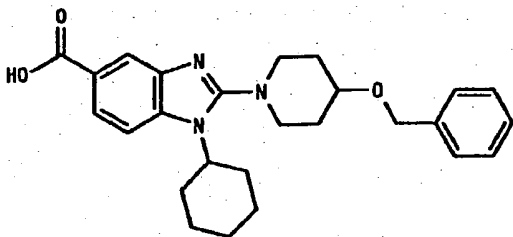
Example No.	93	1H NMR ( $\delta$ ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.28 (1H, d, J=8.9Hz), 8.05 (1H, d, J=8.7Hz), 7.72 (2H, d, J=8.7Hz), 7.38 (4H, d, J=7.2Hz), 7.31 (4H, t, J=7.3Hz), 7.21-7.17 (4H, m), 4.37 (1H, m), 4.26 (1H, t, J=7.9Hz), 4.01 (2H, t, J=6.2Hz), 2.57 (2H, m), 2.50-2.20 (2H, m), 2.10-2.00 (2H, m), 2.00-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.20 (3H, m).
Purity	>90% (NMR)	
MS	531 (M+1)	

Table 22

Example No.	94	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.27 (1H, d, J=9.0Hz), 8.05 (1H, d, J=8.7Hz), 7.75-7.70 (3H, m), 7.56 (1H, d, J=8.4Hz), 7.55-7.35 (6H, m), 7.22 (2H, d, J=8.7Hz), 5.11 (2H, s), 4.36 (1H, m), 2.40-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.20 (3H, m).
Purity	> 90% (NMR)	
MS	537 (M+1)	

Example No.	95	1H NMR (δ) ppm
		300Hz, DMSO-d6 12.9 (1H, brs), 8.02 (1H, s), 7.82 (2H, m), 7.40-7.25 (5H, m), 4.58 (2H, s), 4.09 (1H, m), 3.71 (1H, m), 3.49 (2H, m), 3.21 (2H, m), 2.35-1.30 (14H, m).
Purity	> 90% (NMR)	
MS	434 (M+1)	

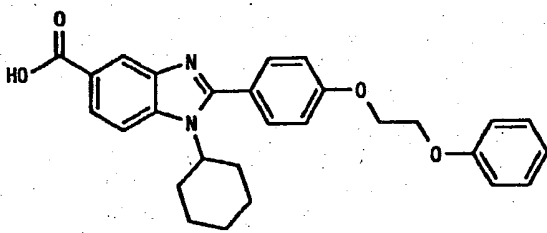
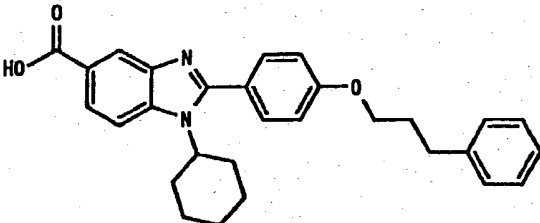
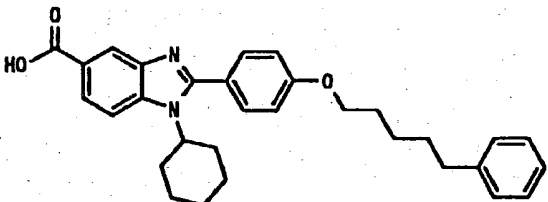
Example No.	96	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, d, J=1.3Hz), 8.27 (1H, d, J=8.8Hz), 8.05 (1H, d, J=8.8Hz), 7.76 (2H, d, J=8.7Hz), 7.40-7.25 (4H, m), 7.06-6.90 (3H, m), 4.53-4.26 (5H, m), 2.40-2.18 (2H, m), 2.12-1.56 (5H, m), 1.50-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	457 (M+1)	

Table 23

Example No.	97	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, d, J=1.3Hz), 8.29 (1H, d, J=8.8Hz), 8.05 (1H, dd, J=8.8, 1.3Hz), 8.42 (2H, d, J=8.8Hz), 7.37-7.16 (7H, m), 4.48-4.30 (1H, m), 4.12 (2H, t, J=6.2Hz), 2.83-2.70 (2H, m), 2.40-1.50 (9H, m), 1.59-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	455 (M+1)	

Example No.	98	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.28 (1H, d, J=1.3Hz), 8.21 (1H, d, J=8.8Hz), 8.01 (1H, d, J=10.1Hz), 7.70 (2H, d, J=8.7Hz), 7.33-7.12 (7H, m), 4.44-4.28 (1H, m), 4.10 (2H, t, J=6.3Hz), 2.62 (2H, t, J=7.4Hz), 2.39-2.15 (2H, m), 2.10-1.18 (14H, m)
Purity	> 90% (NMR)	
MS	483 (M+1)	

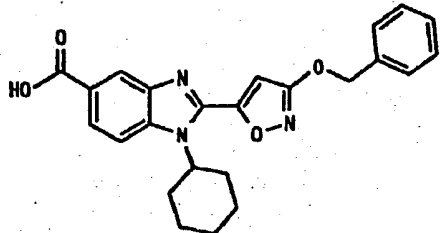
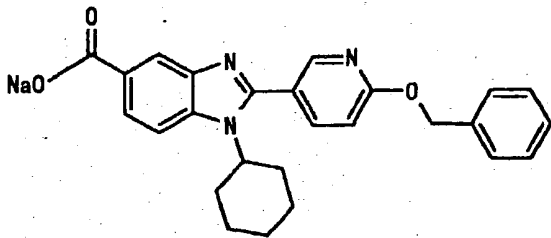
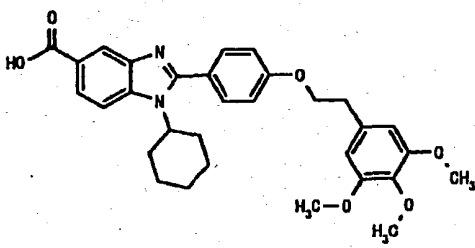
Example No.	99	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.93 (1H, brs), 8.30 (1H, d, J=1.4Hz), 8.04 (1H, d, J=8.7Hz), 7.92 (1H, dd, J=8.7, 1.4Hz), 7.59-7.34 (5H, m), 7.07 (1H, s), 5.38 (2H, s), 4.78-4.60 (1H, m), 2.32-2.14 (2H, m), 2.03-1.28 (8H, m)
Purity	> 90% (NMR)	
MS	418 (M+1)	

Table 24

Example No.	100	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.46 (1H, d, J=2.1Hz), 8.16 (1H, s), 8.00 (1H, dd, J=8.5, 2.1Hz), 7.87 (1H, d, J=8.5Hz), 7.68 (1H, d, J=8.5Hz), 7.55-7.30 (5H, m), 7.08 (1H, d, J=8.5Hz), 5.45 (2H, s), 4.25-4.08 (1H, m), 2.39-2.18 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.45-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	427 (M+1)	

Example No.	101	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.31 (1H, d, J=6.9Hz), 8.06 (1H, d, J=8.4Hz), 7.76 and 7.29 (4H, ABq, J=8.9Hz), 6.68 (2H, s), 4.37 (1H, m), 4.35 (2H, t, J=7.0Hz), 3.79 (6H, s), 3.63 (3H, s), 3.04 (2H, t, J=6.9Hz), 2.30 (2H, m), 2.04 (2H, m), 1.86 (2H, m), 1.65 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	531 (M+1)	

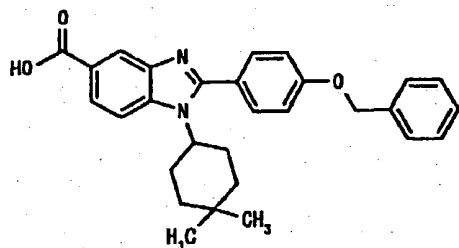
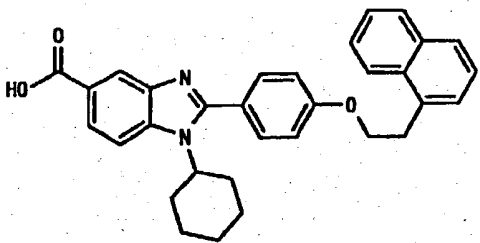
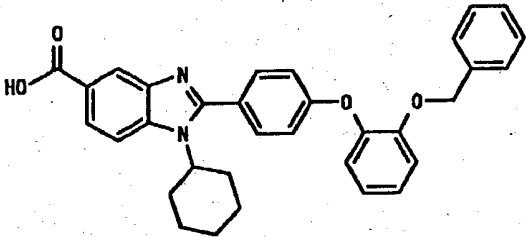
Example No.	102	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.88 (1H, s), 8.34 (1H, s), 7.86 (1H, d, J=8.5Hz), 7.73 (1H, d, J=8.5Hz), 7.63 and 7.23 (4H, ABq, J=8.7Hz), 7.52-7.35 (5H, m), 5.22 (2H, s), 4.31 (1H, m), 2.39 (2H, m), 1.79 (2H, m), 1.53 (2H, m), 1.31 (2H, m), 1.11 (3H, s), 0.95 (3H, s)
Purity	> 90% (NMR)	
MS	455 (M+1)	

Table 25

Example No.	103	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.79 (1H, brs), 8.22 (2H, s), 8.02-7.78 (4H, m), 7.63-7.42 (6H, m), 7.20-7.09 (2H, m), 4.43 (2H, s), 4.27 (1H, brt, J=12.2Hz), 3.59 (2H, s), 2.39-2.15 (2H, m), 1.98-1.72 (4H, m), 1.68-1.59 (1H, m), 1.43-1.12 (3H, m)
Purity	> 90% (NMR)	
MS	491 (M+1)	

Example No.	104	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, s), 8.23 (1H, s), 7.94 and 7.86 (2H, ABq, J=8.6Hz), 7.64 and 7.05 (4H, A'B'q, J=8.7Hz), 7.32-7.09 (9H, m), 5.13 (2H, s), 4.28 (1H, brt, J=12.2Hz), 2.36-2.19 (2H, m), 1.95-1.77 (4H, m), 1.66-1.56 (1H, m), 1.46-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	519 (M+1)	

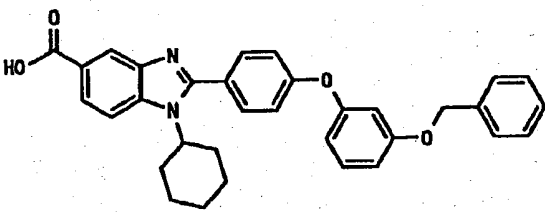
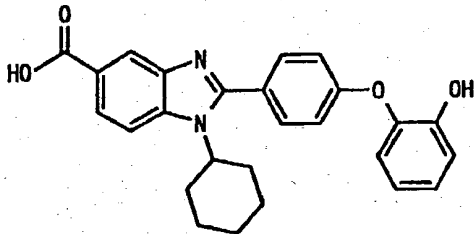
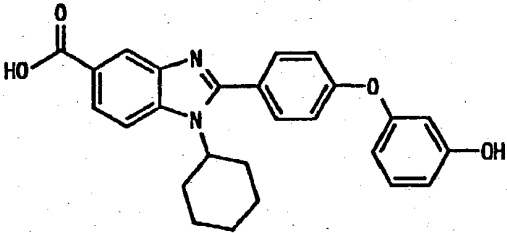
Example No.	105	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.94 and 7.87 (2H, ABq, J=8.6Hz), 7.68 and 7.17 (4H, A'B'q, J=8.7Hz), 7.46-7.33 (6H, m), 6.93 and 6.75 (2H, A''B''q, J=8.2Hz), 6.82 (1H, s), 5.13 (2H, s), 4.30 (1H, brt, J=12.2Hz), 2.39-2.18 (2H, m), 1.98-1.77 (4H, m), 1.71-1.59 (1H, m), 1.48-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	519 (M+1)	

Table 26

Example No.	106	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.89 (1H, brs), 9.73 (1H, s), 8.24 (1H, s), 8.03 and 7.91 ( 2H, ABq, J=8.7Hz), 7.66 and 7. 04 (4H, A' B' q, J=8.7Hz), 7. 16-7.03 (3H, m), 6.89 (2H, t, J=9.2Hz), 4.33 (1H, brt, J=1 2.2Hz), 2.40-2.18 (2H, m), 2. 00-1.78 (4H, m), 1.70-1.58 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	429 (M+1)	

Example No.	107	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.98 (1H, brs), 9.82 (1H, br s), 8.27 (1H, s), 8.09 and 7.9 4 (2H, ABq, J=8.7Hz), 7.74 and 7.22 (4H, A' B' q, J=8.7Hz), 7.28-7.22 (1H, m), 6.67-6.5 4 (3H, m), 4.35 (1H, brt, J=12 .2Hz), 2.40-2.20 (2H, m), 2. 05-1.80 (4H, m), 1.72-1.59 ( 1H, m), 1.50-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	429 (M+1)	

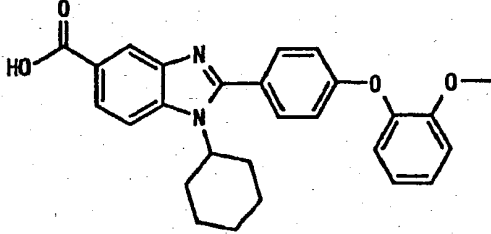
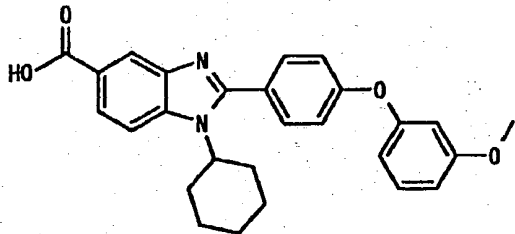
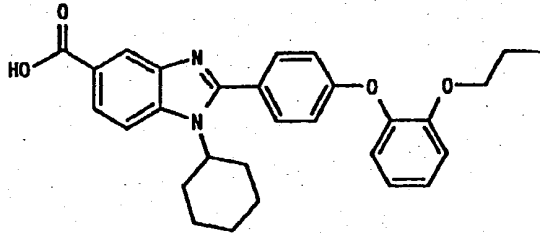
Example No.	108	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, s), 8.01 and 7.90 (2 H, ABq, J=8.7Hz), 7.65 and 7. 03 (4H, A' B' q, J=8.7Hz), 7.3 2-7.20 (3H, m), 7.08-7.03 (1 H, m), 4.32 (1H, brt, J=12.2H z), 3.77 (3H, s), 2.36-2.20 ( 2H, m), 2.00-1.78 (4H, m), 1. 71-1.59 (1H, m), 1.44-1.11 ( 3H, m)
Purity	> 90% (NMR)	
MS	443 (M+1)	

Table 27

Example No.	109	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.75 (1H, s), 8.24 (1H, s), 7.96 and 7.87 (2H, ABq, J=9.0 Hz), 7.69 and 7.19 (4H, A'B'q, J=8.6 Hz), 7.37 (1H, t, J=7.1 Hz), 6.84-6.70 (3H, m), 4.31 (1H, brt, J=12.2 Hz), 3.78 (3H, s), 2.39-2.20 (2H, m), 1.98-1.78 (4H, m), 1.76-1.60 (1H, m), 1.48-1.13 (3H, m)
Purity	> 90% (NMR)	
MS	443 (M+1)	

Example No.	110	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31 (1H, s), 8.26 and 8.04 (2H, ABq, J=8.8 Hz), 7.75 and 7.71 (4H, A'B'q, J=8.8 Hz), 7.32-7.03 (4H, m), 4.34 (1H, brt, J=12.2 Hz), 3.94 (2H, t, J=6.3 Hz), 2.40-2.19 (2H, m), 2.11-1.81 (4H, m), 1.72-1.16 (6H, m), 0.71 (3H, t, J=7.3 Hz)
Purity	> 90% (NMR)	
MS	471 (M+1)	

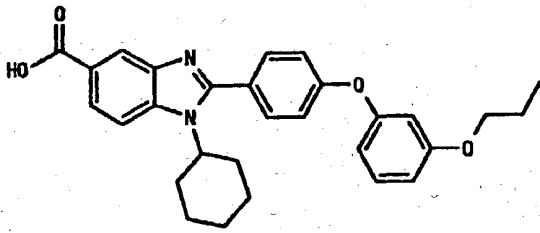
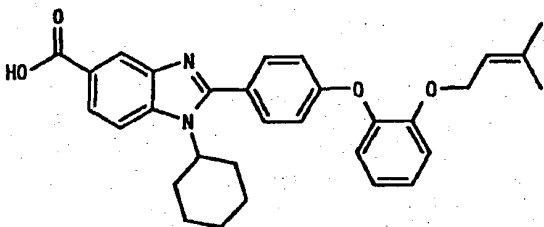
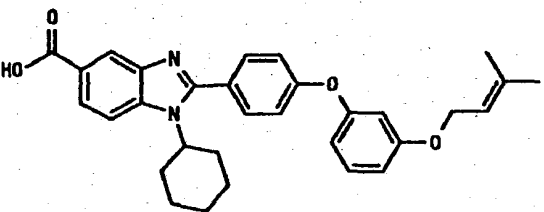
Example No.	111	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.22 (1H, s), 7.91 and 7.87 (2H, ABq, J=8.7 Hz), 7.68 and 7.18 (4H, A'B'q, J=8.7 Hz), 7.35 (1H, t, J=8.5 Hz), 6.80 (1H, d, J=9.0 Hz), 6.72-6.68 (2H, m), 4.30 (1H, brt, J=12.2 Hz), 3.94 (2H, t, J=6.5 Hz), 2.39-2.18 (2H, m), 1.97-1.58 (7H, m), 1.45-1.20 (3H, m), 0.97 (3H, t, J=7.4 Hz)
Purity	> 90% (NMR)	
MS	471 (M+1)	

Table 28

Example No.	112	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.73 (1H, s), 8.22 (1H, s), 7.94 and 7.85 (2H, ABq, J=9.3 Hz), 7.61 and 7.01 (4H, A'B'q, J=8.6 Hz), 7.25-7.00 (4H, m), 5.25 (2H, brs), 4.55 (2H, d, J=6.6 Hz), 4.29 (1H, brt, J=12.2 Hz), 2.38-2.18 (2H, m), 1.96-1.78 (4H, m), 1.70-1.56 (1H, m), 1.67 (3H, s), 1.60 (3H, s), 1.48-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	497 (M+1)	

Example No.	113	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, s), 8.23 (1H, s), 7.95 and 7.86 (2H, ABq, J=8.9 Hz), 7.69 and 7.18 (4H, A'B'q, J=8.9 Hz), 7.35 (1H, t, J=8.3 Hz), 6.81-6.69 (3H, m), 5.41 (2H, brs), 4.54 (2H, d, J=6.6 Hz), 4.31 (1H, brt, J=12.2 Hz), 2.41-2.18 (2H, m), 1.98-1.76 (4H, m), 1.73 (3H, s), 1.70-1.58 (1H, m), 1.68 (3H, s), 1.45-1.17 (3H, m)
Purity	> 90% (NMR)	
MS	497 (M+1)	

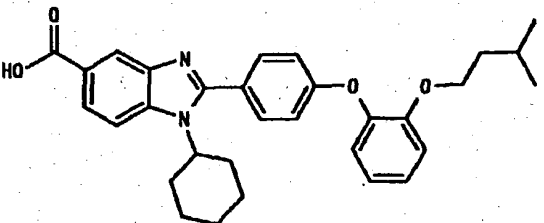
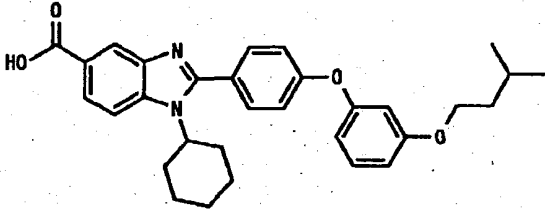
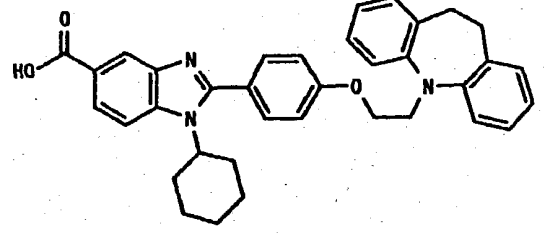
Example No.	114	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.73 (1H, s), 8.22 (1H, s), 7.94 and 7.85 (2H, ABq, J=8.4 Hz), 7.60 and 6.99 (4H, A'B'q, J=8.6 Hz), 7.29-7.00 (4H, m), 4.29 (1H, brt, J=12.2 Hz), 3.99 (2H, t, J=6.3 Hz), 2.41-2.20 (2H, m), 1.95-1.76 (4H, m), 1.70-1.14 (7H, m), 0.76 (3H, d, J=6.6 Hz)
Purity	> 90% (NMR)	
MS	499 (M+1)	



Table 29

Example No.	115	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.93 and 7.87 (2H, ABq, J=8.6Hz), 7.69 and 7.19 (4H, A'B'q, J=8.6Hz), 7.35 (1H, t, J=7.8Hz), 6.82-6.69 (3H, m), 4.30 (1H, brt, J=12.2Hz), 4.00 (2H, t, J=6.9Hz), 2.38-2.20 (2H, m), 1.97-1.54 (8H, m), 1.47-1.20 (3H, m), 0.93 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	499 (M+1)	

Example No.	116	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.25 (1H, d, J=8.9Hz), 8.03 (1H, d, J=8.8Hz), 7.68 (2H, d, J=8.8Hz), 7.24 (2H, d, J=7.2Hz), 7.19-7.10 (6H, m), 6.94 (2H, t, J=7.2Hz), 4.34 (1H, m), 4.19 (4H, brs), 3.10 (4H, brs), 2.40-2.15 (2H, m), 2.10-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.20 (3H, m).
Purity	> 90% (NMR)	
MS	557 (M+1)	

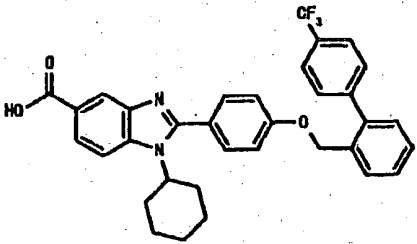
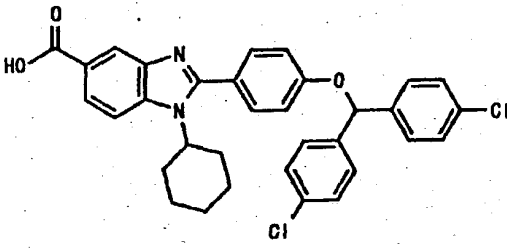
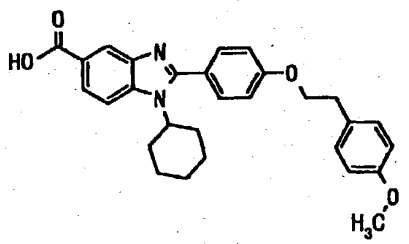
Example No.	117	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.8 (1H, brs), 8.22 (1H, s), 7.98 (1H, d, J=8.7Hz), 7.87 (1H, d, J=8.6Hz), 7.80 (2H, d, J=8.2Hz), 7.72-7.67 (3H, m), 7.59 (2H, d, J=8.7Hz), 7.54-7.51 (2H, m), 7.42-7.41 (1H, m), 7.11 (2H, d, J=8.8Hz), 5.09 (2H, s), 4.27 (1H, m), 2.40-2.15 (2H, m), 2.00-1.75 (4H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	571 (M+1)	

Table 30

Example No.	118	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.3 (1H, brs), 8.30 (1H, s), 8.25 (1H, d, J=8.9Hz), 8.04 (1H, d, J=8.7Hz), 7.72 (2H, d, J=8.8Hz), 7.57 (4H, d, J=8.6Hz), 7.47 (4H, d, J=8.6Hz), 7.33 (2H, d, J=8.9Hz), 6.84 (1H, s), 4.33 (1H, m), 2.45-2.10 (2H, m), 2.10-1.95 (2H, m), 1.95-1.70 (2H, m), 1.70-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	571 (M+1)	

Example No.	119	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32-8.30 (2H, m), 8.07-8.03 (1H, m), 7.74 and 6.90 (4H, ABq, J=8.7Hz), 4.37 (1H, m), 4.31 (2H, t, J=6.8Hz), 3.74 (3H, s), 3.04 (2H, t, J=6.7Hz), 2.30 (2H, m), 2.02 (2H, m), 1.86 (2H, m), 1.63 (1H, m), 1.55-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	471 (M+1)	

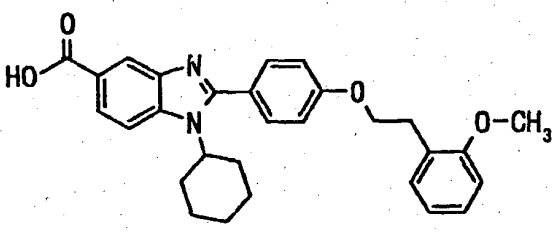
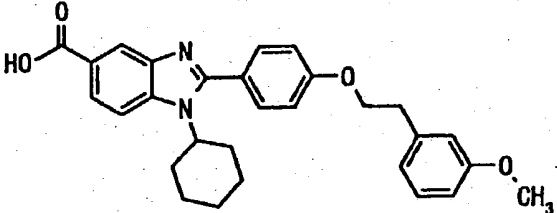
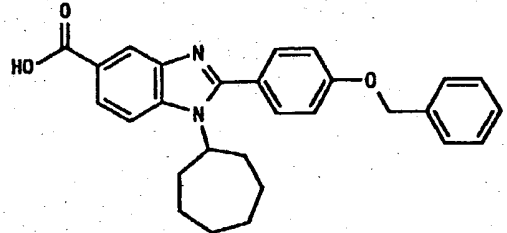
Example No.	120	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.99 (1H, d, J=8.7Hz), 7.88 (1H, d, J=8.4Hz), 7.61 and 7.16 (4H, ABq, J=8.6Hz), 7.30-7.22 (2H, m), 7.01 (2H, d, J=8.1Hz), 6.92 (1H, t, J=7.5Hz), 4.28 (1H, m), 4.25 (2H, t, J=7.2Hz), 3.83 (3H, s), 3.07 (2H, t, J=7.1Hz), 2.28 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	471 (M+1)	

Table 31

Example No.	121	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.85 (1H, brs), 8.24 (1H, s), 8.01 (1H, d, J=8.7Hz), 7.90 (1H, d, J=8.6Hz), 7.62 and 7 .17 (4H, ABq, J=8.7Hz), 7.24 (1H, m), 6.94 (2H, m), 6.82 (1 H, m), 4.32 (2H, t, J=6.7Hz), 3.76 (3H, s), 3.07 (2H, t, J=6 .7Hz), 2.29 (2H, m), 2.00-1. 75 (4H, m), 1.70-1.55 (1H, m) , 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	471 (M+1)	

Example No.	122	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.8 (1H, brs), 8.22 (1H, s), 7.87 (2H, m), 7.62 (2H, d, J=8 .1Hz), 7.60-7.20 (7H, m), 5. 23 (2H, s), 4.46 (1H, m), 2.50 -2.30 (2H, m), 1.70-1.40 (10 H, m).
Purity	> 90% (NMR)	
MS	441 (M+1)	

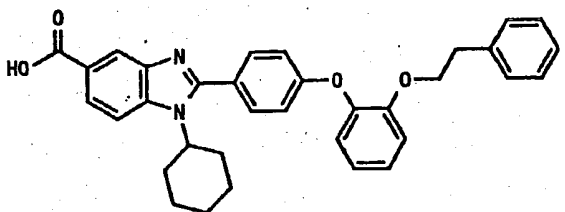
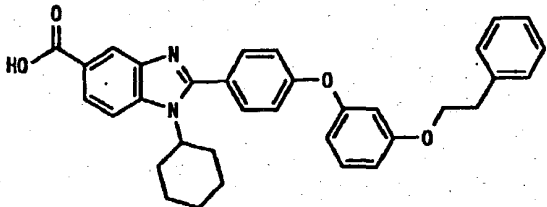
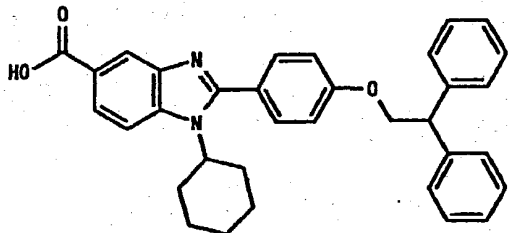
Example No.	123	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, s), 7.97 (1H, d, J=9 .0Hz), 7.87 (1H, d, J=8.4Hz) , 7.65 (2H, d, J=8.7Hz), 7.40 -7.05 (9H, m), 7.03 (2H, d, J= 8.4Hz), 4.31 (1H, m), 4.18 (2 H, t, J=6.6Hz), 2.81 (2H, t, J =6.3Hz), 2.40-2.20 (2H, m), 2.00-1.70 (4H, m), 1.70-1.5 0 (1H, m), 1.50-1.05 (3H, m).
Purity	> 90% (NMR)	
MS	533 (M+1)	

Table 32

Example No.	124	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.1 (1H, brs), 8.29 (1H, s), 8.17 (1H, d, J=8.7Hz), 7.99 (1H, d, J=8.7Hz), 7.77 (2H, d, J=8.7Hz), 7.40-7.20 (8H, m), 6.84 (1H, d, J=9.3Hz), 6.75-6.72 (2H, m), 4.36 (1H, m), 4.22 (2H, t, J=6.8Hz), 3.04 (2H, t, J=6.7Hz), 2.40-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	533 (M+1)	

Example No.	125	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.28 (1H, d, J=8.7Hz), 8.05 (1H, d, J=9.0Hz), 7.73 (2H, d, J=9.0Hz), 7.43 (4H, d, J=7.2Hz), 7.36-7.20 (8H, m), 4.74 (2H, d, J=7.5Hz), 4.57 (1H, t, J=7.5Hz), 4.38 (1H, m), 2.40-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.85 (2H, m), 1.85-1.55 (1H, m), 1.55-1.20 (3H, m).
Purity	> 90% (NMR)	
MS	517 (M+1)	

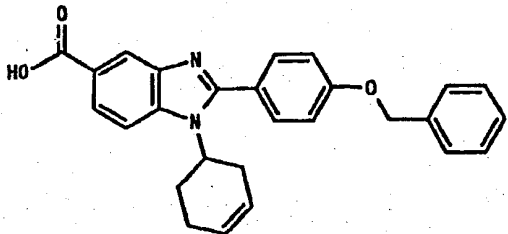
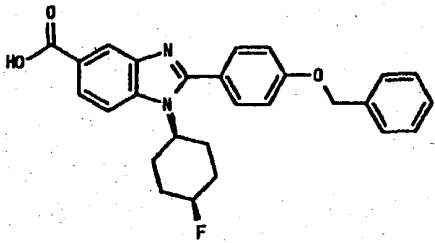
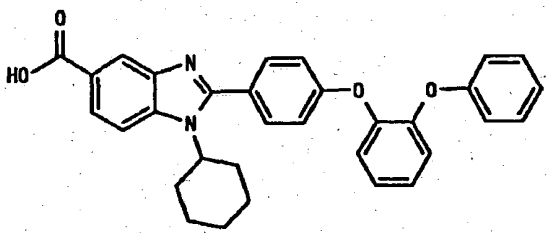
Example No.	126	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.14 (1H, d, J=8.7Hz), 8.03 (1H, d, J=8.7Hz), 7.77 (2H, d, J=9.0Hz), 7.52-7.31 (7H, m), 5.74 (2H, m), 5.26 (2H, s), 4.61 (1H, m), 2.96 (1H, m), 2.60-2.10 (5H, m).
Purity	> 90% (NMR)	
MS	425 (M+1)	

Table 33

Example No.	127	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.2 (1H, brs), 8.33 (1H, s), 8.12 (1H, d, J=8.7Hz), 7.96 (1H, d, J=8.8Hz), 7.79 (2H, d, J=8.7Hz), 7.52-7.32 (7H, m), 5.26 (2H, s), 4.92 (1H, d, J=49.4Hz), 4.57 (1H, m), 2.65-2.35 (2H, m), 2.25-1.50 (6H, m).
Purity	> 90% (NMR)	
MS	445 (M+1)	

Example No.	128	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 7.92 and 7.85 (2H, ABq, J=8.6Hz), 7.61 and 7.06 (4H, A'B'q, J=8.6Hz), 7.36-6.91 (9H, m), 4.24 (1H, brt, J=12.2Hz), 2.35-2.15 (2H, m), 1.95-1.75 (4H, m), 1.70-1.58 (1H, m), 1.48-1.14 (3H, m)
Purity	> 90% (NMR)	
MS	505 (M+1)	

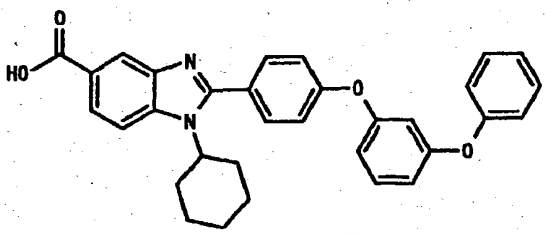
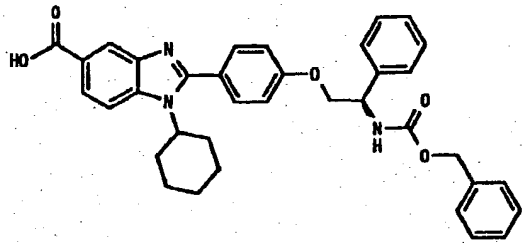
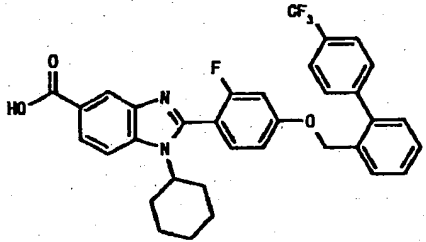
Example No.	129	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 7.92 and 7.86 (2H, ABq, J=8.6Hz), 7.69 and 7.22 (4H, A'B'q, J=8.6Hz), 7.52-7.39 (1H, m), 7.47 and 7.41 (2H, A''B''q, J=8.1Hz), 6.91 (1H, d, J=8.0Hz), 6.89 (1H, d, J=8.2Hz), 6.75 (1H, s), 4.36-4.18 (1H, m), 2.38-2.17 (2H, m), 1.95-1.76 (4H, m), 1.70-1.59 (1H, m), 1.44-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	505 (M+1)	

Table 34

Example No.	130	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 7.69 (2H, d, J=8.6Hz), 7.49-7.21 (11H, m), 5.08 and 5.03 (2H, ABq, J=12.6Hz), 5.07-4.99 (1H, m), 4.26 (2H, d, J=6.6Hz), 2.40-2.18 (2H, m), 2.04-1.77 (4H, m), 1.70-1.58 (1H, m), 1.48-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	590 (M+1)	

Example No.	131	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.11 (1H, d, J=9.0Hz), 7.96 (1H, d, J=8.4Hz), 7.80 (2H, d, J=8.1Hz), 7.72-7.41 (7H, m), 7.12 (1H, d, J=12.6Hz), 7.01 (1H, d, J=8.4Hz), 5.12 (2H, s), 4.06 (1H, m), 2.35-2.10 (2H, m), 2.00-1.75 (4H, m), 1.75-1.55 (1H, m), 1.60-1.20 (3H, m).
Purity	> 90% (NMR)	
MS	589 (M+1)	

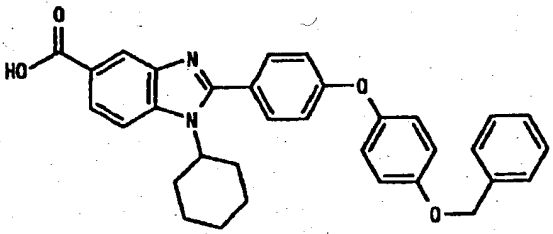
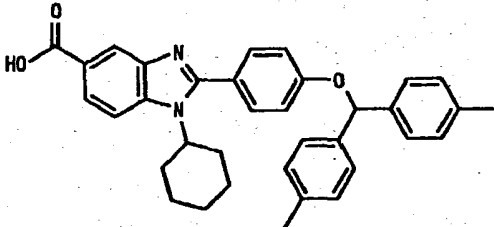
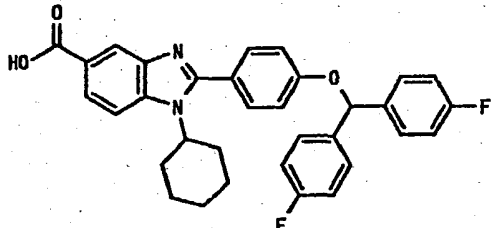
Example No.	132	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.8 (1H, brs), 8.23 (1H, s), 7.97 (1H, d, J=8.7Hz), 7.87 (1H, d, J=8.6Hz), 7.66 (2H, d, J=8.6Hz), 7.49-7.33 (5H, m), 7.17-7.05 (6H, m), 5.12 (2H, s), 4.31 (1H, m), 2.40-2.15 (2H, m), 2.05-1.20 (8H, m).
Purity	> 90% (NMR)	
MS	519 (M+1)	

Table 35

Example No.	133	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.57(1H, s), 8.01(1H, d, J=8.7Hz), 7.66(1H, d, J=8.7Hz), 7.51(2H, d, J=8.7Hz), 7.31(4H, d, J=8.0Hz), 7.16(4H, d, J=8.0Hz), 7.09(2H, d, J=8.7Hz), 6.26(1H, s), 4.37(1H, m), 2.41-2.28(2H, m), 2.33(6H, s), 2.03-1.84(4H, m), 1.77(1H, m), 1.45-1.20(3H, m)
Purity	> 90% (NMR)	
MS	531(M+1)	

Example No.	134	1H NMR(δ) ppm
		8.59(1H, d, J=1.5Hz), 8.02(1H, dd, J=8.7, 1.5Hz), 7.68(1H, d, J=8.7Hz), 7.54(2H, d, J=8.8Hz), 7.39(4H, dd, J=8.7, 5.3Hz), 7.08(4H, d, J=8.7Hz), 7.05(2H, d, J=8.8Hz), 6.29(1H, s), 4.36(1H, m), 2.43-2.19(2H, m), 2.04-1.85(4H, m), 1.78(1H, m), 1.45-1.23(3H, m).
Purity	> 90% (NMR)	
MS	539(M+1)	

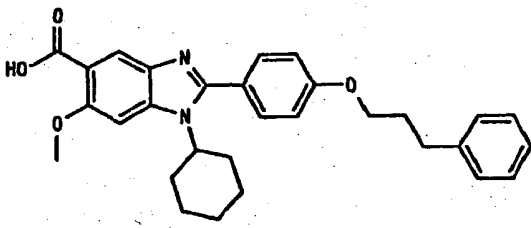
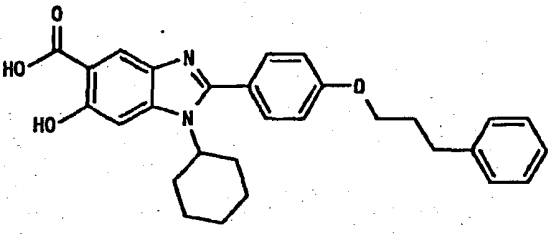
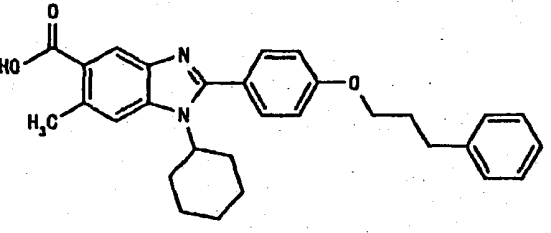
Example No.	135	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.34(1H, brs), 7.93(1H, s), 7.55(1H, d, J=8.6Hz), 7.33-7.15(6H, m), 7.11(2H, d, J=8.6Hz), 4.30-4.20(1H, m), 4.07(2H, t, J=6.3Hz), 3.93(3H, s), 2.78(2H, t, J=7.4Hz), 2.35-2.19(2H, m), 2.12-2.00(2H, m), 1.91-1.79(4H, m), 1.69-1.60(1H, m), 1.47-1.20(3H, m)
Purity	> 90% (NMR)	
MS	485(M+1)	

Table 36

Example No.	136	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.13 (1H, s), 7.65 (2H, d, J=8.7Hz), 7.63 (1H, s), 7.35-7.12 (7H, m), 4.35-4.20 (1H, m), 4.10 (1H, t, J=6.3Hz), 2.78 (2H, t, J=7.5Hz), 2.33-1.78 (8H, m), 1.70-1.16 (4H, m)
Purity	> 90% (NMR)	
MS	471 (M+1)	

Example No.	137	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, s), 8.11 (1H, s), 7.76 (2H, d, J=9.0Hz), 7.37-7.16 (7H, m), 4.43-4.30 (1H, m), 4.13 (2H, t, J=6.3Hz), 2.84-2.68 (5H, m), 2.42-2.22 (2H, m), 2.18-1.80 (6H, m), 1.70-1.20 (4H, m)
Purity	> 90% (NMR)	
MS	469 (M+1)	

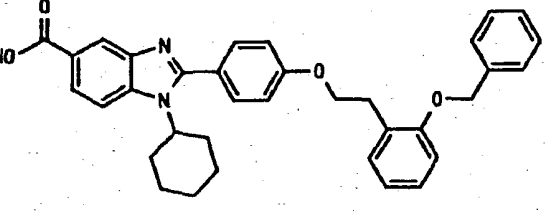
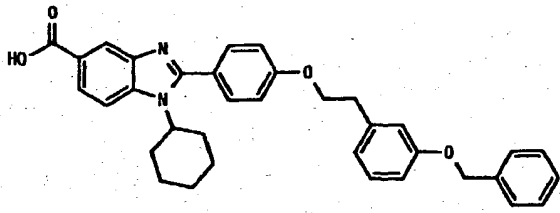
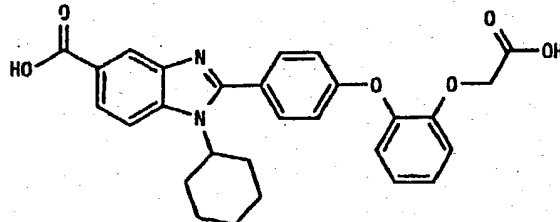
Example No.	138	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.73 (1H, brs), 8.22 (1H, s), 7.76 (1H, d, J=8.7Hz), 7.85 (1H, d, J=8.7Hz), 7.54-7.49 (4H, m), 7.42-7.21 (5H, m), 7.11-7.09 (3H, m), 6.93 (1H, m), 5.17 (2H, s), 4.29 (3H, m), 3.11 (2H, m), 2.40-2.20 (2H, m), 1.99-1.23 (8H, m)
Purity	> 90% (NMR)	
MS	547 (M+1)	



Table 37

Example No.	139	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.73 (1H, brs), 8.22 (1H, s), 7.93 (1H, d, J=8.7Hz), 7.73 (1H, m), 7.60-7.57 (2H, m), 7 .47-6.90 (1H, m), 5.11 (2H, s ), 4.33-4.28 (3H, m), 3.09-3 .04 (2H, t, J=6.7Hz), 2.35-2 .20 (2H, m), 1.95-1.10 (8H, m )
Purity	> 90% (NMR)	
MS	547 (M+1)	

Example No.	140	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.83 (2H, brs), 8.22 (1H, s) , 7.94 (1H, d, J=8.7Hz), 7.85 (1H, d, J=8.4Hz), 7.63-7.60 (2H, m), 7.26-7.03 (6H, m), 4 .73 (2H, s), 4.30 (1H, m), 2.4 0-2.15 (2H, m), 2.00-1.20 (8 H, m)
Purity	> 90% (NMR)	
MS	487 (M+1)	

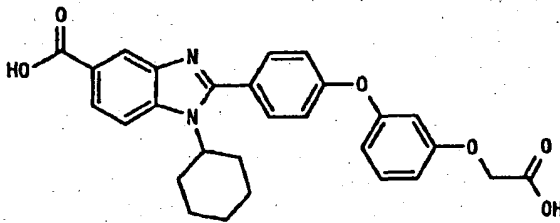
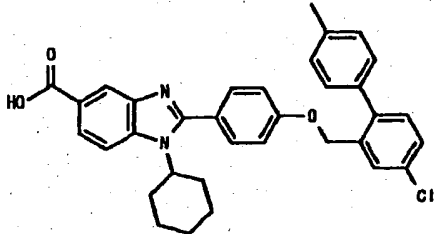
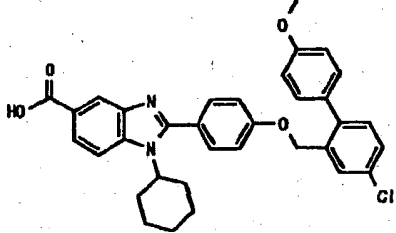
Example No.	141	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.87 (1H, brs), 8.24 (1H, s) , 7.97 (1H, d, J=9.0Hz), 7.87 (1H, d, J=8.7Hz), 7.69 and 7. 19 (4H, ABq, J=8.7Hz), 7.36 (1H, t, J=8.7Hz), 6.80-6.72 (3H, m), 4.71 (2H, s), 4.32 (1H, m), 2.29 (2H, m), 1.95-1.25 (8H, m)
Purity	> 90% (NMR)	
MS	487 (M+1)	

Table 38

Example No.	142	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.27 (1H, d, J=8.7Hz), 8.05 (1H, d, J=9.0Hz), 7.76-7.72 (3H, m), 7.54 (1H, d, J=8.4Hz), 7.39-7.22 (7H, m), 5.11 (1H, s), 4.36 (1H, m), 2.35 (3H, s), 2.35-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	551 (M+1)	

Example No.	143	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.1 (1H, brs), 8.30 (1H, s), 8.24 (1H, d, J=8.8Hz), 8.03 (1H, d, J=8.7Hz), 7.74-7.71 (3H, m), 7.52 (1H, d, J=8.3Hz), 7.40-7.36 (3H, m), 7.23 (2H, d, J=8.8Hz), 7.01 (2H, d, J=8.7Hz), 5.11 (2H, s), 4.35 (1H, m), 3.79 (3H, s), 2.45-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	567 (M+1)	

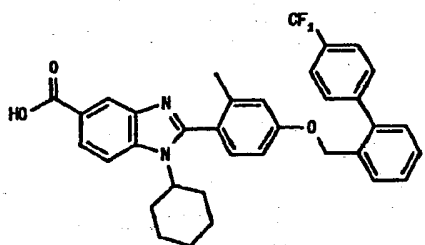
Example No.	144	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.0 (1H, brs), 8.31 (1H, s), 8.23 (1H, d, J=8.7Hz), 8.04 (1H, d, J=8.7Hz), 7.80 (2H, d, J=8.3Hz), 7.70-7.66 (3H, m), 7.55-7.40 (4H, m), 7.03-6.95 (2H, m), 5.08 (2H, s), 4.03 (1H, m), 2.40-2.15 (2H, m), 2.18 (3H, s), 2.05-1.70 (4H, m), 1.70-1.50 (1H, m), 1.50-1.10 (3H, m).
Purity	> 90% (NMR)	
MS	585 (M+1)	

Table 39

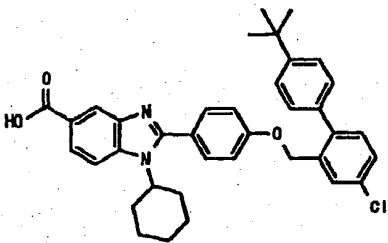
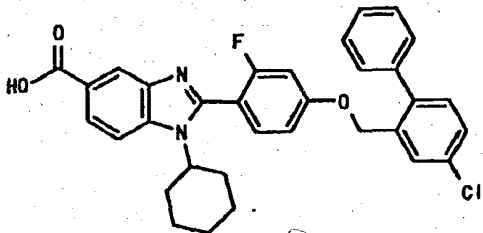
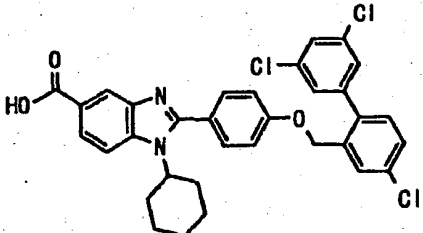
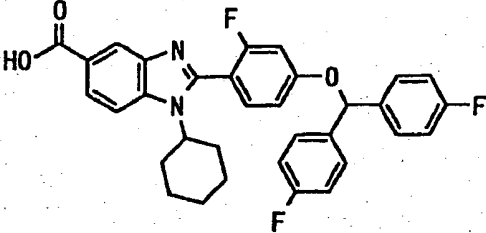
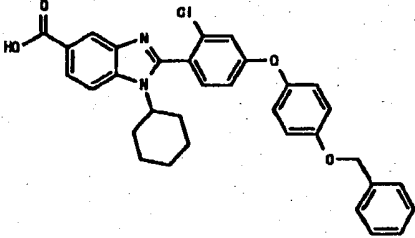
Example No.	145	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.23 (1H, d, J=8.8Hz), 8.02 (1H, d, J=8.7Hz), 7.73-7.71 (3H, m), 7.54 (1H, d, J=8.3Hz), 7.48 (2H, d, J=8.4Hz), 7.41-7.37 (3H, m), 7.22 (2H, d, J=8.7Hz), 5.13 (2H, s), 4.34 (1H, m), 2.40-2.20 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m), 1.31 (9H, s).
Purity	> 90% (NMR)	
MS	593 (M+1)	
Example No.	146	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.13 (1H, d, J=8.7Hz), 7.97 (1H, d, J=8.6Hz), 7.76 (1H, d, J=2.1Hz), 7.63 (1H, t, J=8.5Hz), 7.57 (1H, d, J=8.2, 2.2Hz), 7.55-7.35 (6H, m), 7.15 (1H, d, J=12.1Hz), 7.02 (1H, d, J=8.6Hz), 5.10 (2H, s), 4.07 (1H, m), 2.35-2.10 (2H, m), 2.00-1.70 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	555 (M+1)	
Example No.	147	1H NMR (δ) ppm
		300MHz, CDCl3 8.61 (1H, s), 8.04 (1H, d, J=8.7Hz), 7.69 (1H, d, J=8.7Hz), 7.66 (1H, d, J=2.4Hz), 7.59 (2H, d, J=8.7Hz), 7.42 (1H, d, J=8.0, 2.4Hz), 7.38 (1H, t, J=1.8Hz), 7.28 (2H, d, J=1.8Hz), 7.26 (1H, d, J=8.0Hz), 7.03 (2H, d, J=8.7Hz), 4.94 (2H, s), 4.37 (1H, m), 2.43-2.21 (2H, m), 2.17-1.86 (4H, m), 1.79 (1H, m), 1.43-1.26 (3H, m).
Purity	> 90% (NMR)	
MS	605 (M+1)	

Table 40

Example No.	148	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (s, 1H), 7.89 (1H, d, J=8.7Hz), 7.87 (1H, d, J=8.7Hz), 7.63-7.46 (5H, m), 7.30-7.12 (5H, m), 7.08 (1H, d, J=11.0Hz), 6.81 (1H, s), 3.92 (1H, m), 2.15-2.06 (2H, m), 1.89-1.72 (4H, m), 1.61 (1H, m), 1.42-1.09 (3H, m).
Purity	> 90% (NMR)	
MS	557 (M+1)	

Example No.	149	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, d, J=1.5Hz), 7.96 (1H, d, J=9.0Hz), 7.88 (1H, dd, J=9.0, 1.5Hz), 7.58 (1H, d, J=8.7Hz), 7.50-7.30 (5H, m), 7.22-7.00 (6H, m), 5.13 (2H, s), 3.98-3.80 (1H, s), 2.36-1.10 (10H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

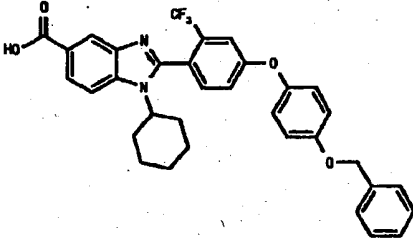
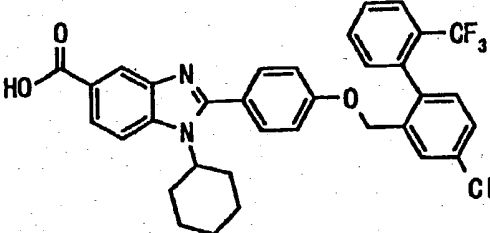
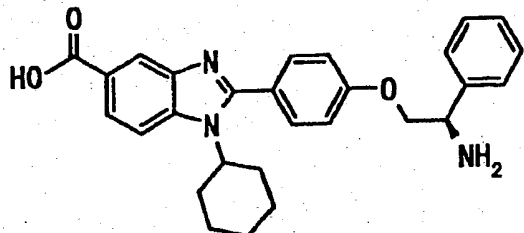
Example No.	150	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 8.95 (1H, d, J=8.4Hz), 7.88 (1H, d, J=8.7Hz), 7.66 (1H, d, J=8.4Hz), 7.52-7.28 (7H, m), 7.23 (2H, d, J=9.3Hz), 7.14 (2H, d, J=8.7Hz), 5.14 (2H, s), 3.90-3.72 (1H, m), 2.20-1.10 (10H, m)
Purity	> 90% (NMR)	
MS	587 (M+1)	

Table 41

Example No.	151	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.18 (1H, s), 7.92-7.78 (3H, m), 7.78-7.58 (3H, m), 7.58-7.44 (4H, m), 7.29 (1H, d, J=8.2Hz), 7.01 (2H, d, J=8.7Hz), 4.88 (1H, d, J=11.8Hz), 4.80 (1H, d, J=11.8Hz), 4.22 (1H, m), 2.37-2.16 (2H, m), 1.95-1.75 (4H, m), 1.64 (1H, m), 1.48-1.14 (3H, m).
Purity	> 90% (NMR)	
MS	605 (M+1)	

Example No.	152	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (2H, m), 7.99-7.80 (2H, m), 7.63-7.08 (9H, m), 4.20-3.98 (4H, m), 2.20-2.15 (2H, m), 1.95-1.74 (4H, m), 1.70-1.54 (1H, m), 1.44-1.14 (3H, m)
Purity	> 90% (NMR)	
MS	456 (M+1)	

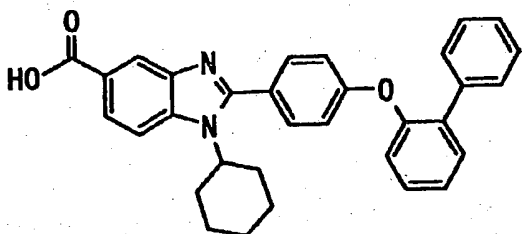
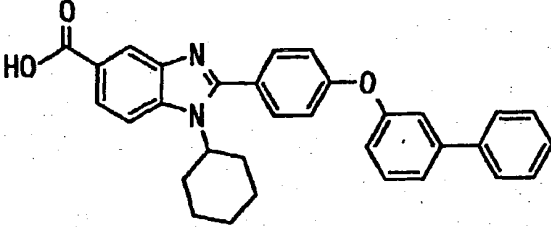
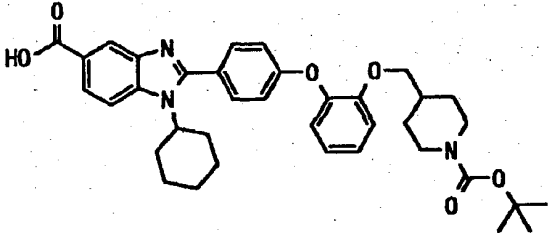
Example No.	153	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.20 (1H, s), 8.93 and 7.83 (2H, ABq, J=8.7Hz), 7.86-7.21 (11H, m), 7.03 (2H, d, J=8.7Hz), 4.20 (1H, brt, J=12.2Hz), 2.32-2.13 (2H, m), 1.92-1.74 (4H, m), 1.69-1.58 (1H, m), 1.45-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	489 (M+1)	

Table 42

Example No.	154	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.94 and 7.86 (2H, ABq, J=8.6Hz), 7.72-7.16 (13H, m), 5.25 (2H, brs), 4.55 (2H, d, J=6.6Hz), 4.31 (1H, brt, J=12.2Hz), 2.37-2.18 (2H, m), 1.98-1.77 (4H, m), 1.70-1.58 (1H, m), 1.48-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	489 (M+1)	

Example No.	155	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 7.85 and 7.61 (2H, ABq, J=8.7Hz), 7.61 and 6.99 (4H, A'B'q, J=8.7Hz), 7.28-7.18 (1H, m), 7.25 (2H, d, J=7.5Hz), 7.07-6.99 (1H, m), 4.30 (1H, brt, J=12.2Hz), 3.83 (2H, d, J=6.0Hz), 3.82-3.72 (1H, m), 2.68-2.49 (2H, m), 2.39-2.21 (2H, m), 1.95-1.80 (4H, m), 1.79-1.60 (2H, m), 1.46-1.22 (5H, m), 1.30 (9H, s), 1.00-0.82 (2H, m)
Purity	> 90% (NMR)	
MS	626 (M+1)	

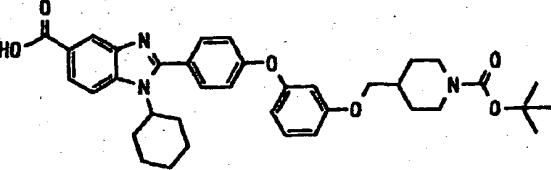
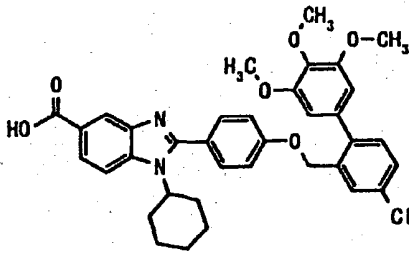
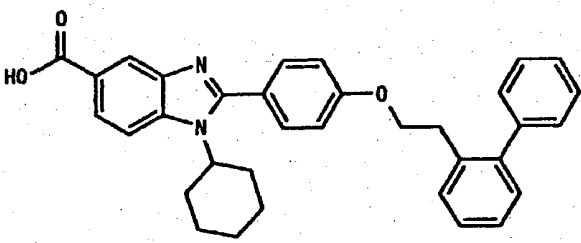
Example No.	156	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.92 and 7.86 (2H, ABq, J=8.7Hz), 7.68 and 7.18 (4H, A'B'q, J=8.7Hz), 7.35 (1H, t, J=8.5Hz), 6.80 (1H, d, J=8.3Hz), 6.72-6.70 (2H, m), 4.30 (1H, brt, J=12.2Hz), 3.99 (2H, brd, J=12.0Hz), 3.85 (2H, d, J=6.3Hz), 2.82-2.62 (2H, m), 2.38-2.20 (2H, m), 1.99-1.59 (8H, m), 1.42-1.03 (5H, m), 1.39 (9H, s)
Purity	> 90% (NMR)	
MS	626 (M+1)	

Table 43

Example No.	157	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.78 (1H, brs), 8.22 (1H, s), 7.96 (1H, d, J=8.6Hz), 7.86 (1H, d, J=8.6Hz), 7.75 (1H, d, J=2.2Hz), 7.60 (2H, d, J=8.4Hz), 7.55 (1H, dd, J=8.3, 2.2Hz), 7.48 (1H, d, J=8.3Hz), 7.18 (2H, d, J=8.4Hz), 6.73 (2H, s), 5.08 (2H, s), 4.23 (1H, m), 3.68 (9H, s), 2.37-2.17 (2H, m), 1.99-1.79 (4H, m), 1.65 (1H, s), 1.49-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	627 (M+1)	

Example No.	158	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, brs), 8.22 (1H, s), 7.93 (2H, d, J=8.7Hz), 7.85 (2H, d, J=8.5Hz), 7.53-7.21 (10H, m), 6.94 (2H, d, J=8.7Hz), 4.30-4.12 (3H, m), 3.05 (2H, m), 2.35-2.15 (2H, m), 1.95-1.75 (4H, m), 1.75-1.55 (1H, m), 1.50-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	517 (M+1)	

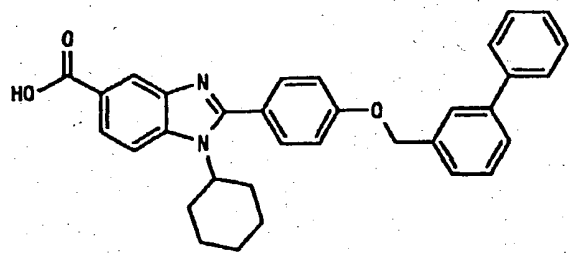
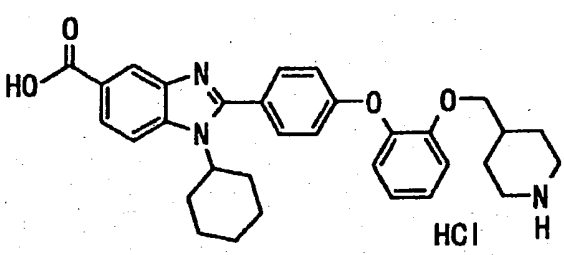
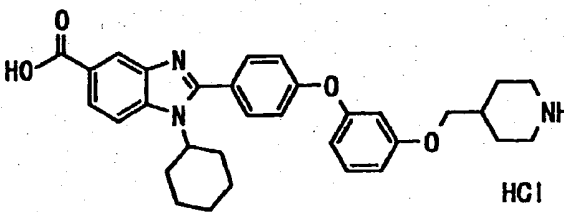
Example No.	159	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.77 (1H, brs), 8.22 (1H, s), 7.95 (1H, d, 8.6Hz), 7.86 (1H, d, 8.6Hz), 7.80 (1H, s), 7.70-7.35 (10H, m), 7.27 (2H, d, J=8.7Hz), 5.30 (2H, s), 4.28 (1H, m), 2.35-2.15 (2H, m), 1.95-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	503 (M+1)	

Table 44

Example No.	160	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.90(1H, brs), 8.59(1h, brs), 8.33(1H, s), 8.18and8.00 (2H, ABq, J=8.5Hz), 7.73and 7.10(4H, A' B' q, J=8.5Hz), 7.32-7.05(4H, m), 4.35(1H, brt, J=12.2Hz), 3.86(2H, d, J=6.3Hz), 3.25-3.08(2H, m), 2.85-2.66(2H, m), 2.40-2.28(2H, m), 2.07-1.14(15H, m)
Purity	> 90% (NMR)	
MS	526 (M+1)	

Example No.	161	1H NMR(δ) ppm
		300MHz, DMSO-d6 9.05(1H, brs), 8.76(1h, brs), 8.31(1H, s), 8.19and8.00 (2H, ABq, J=8.3Hz), 7.79and 7.25(4H, A' B' q, J=8.3Hz), 7.39(1H, brs), 6.86-6.74(4H, m), 4.37(1H, brt, J=12.2Hz), 3.89(2H, d, J=5.0Hz), 3.35-3.18(2H, m), 2.98-2.75(2H, m), 2.38-2.17(2H, m), 2.16-1.15(15H, m)
Purity	> 90% (NMR)	
MS	526 (M+1)	

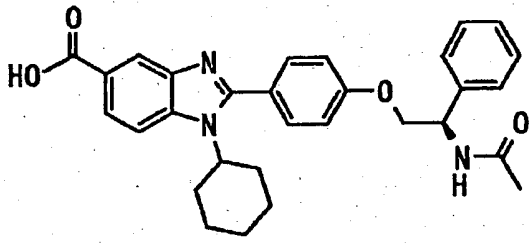
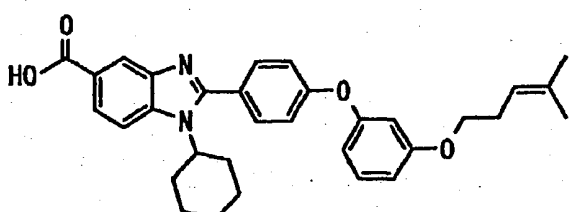
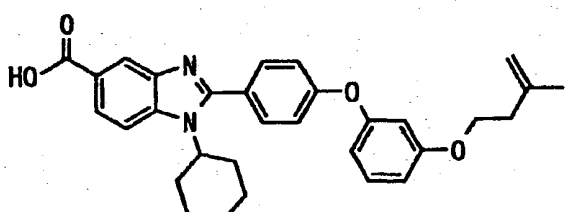
Example No.	162	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.87(1H, brs), 8.58(1H, d, J=6.0Hz), 8.23(1H, s), 7.99and7.80(2H, ABq, J=8.6Hz), 7.61and7.18(4H, A' B' q, J=8.0Hz), 7.45-7.30(5H, m), 5.29(1H, brs), 4.26(1H, brt, J=12.2Hz), 2.37-2.11(2H, m), 2.00-1.71(4H, m), 1.92(3H, s), 1.70-1.52(1H, m), 1.45-1.11(3H, m)
Purity	> 90% (NMR)	
MS	498 (M+1)	



Table 45

Example No.	163	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.95 and 7.86 (2H, ABq, J=8.6Hz), 7.69 and 7.18 (4H, A'B'q, J=8.6Hz), 7.35 (1H, t, J=8.6Hz), 6.80 (1H, d, J=7.5Hz), 6.72-6.69 (2H, m), 5.20 (1H, t, J=3.7Hz), 4.31 (1H, brt, J=12.2Hz), 3.95 (2H, t, J=6.8Hz), 2.49-2.19 (4H, m), 1.97-1.76 (4H, m), 1.68 (3H, s), 1.67-1.54 (1H, m), 1.61 (3H, s), 1.45-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	511 (M+1)	

Example No.	164	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.20 (1H, s), 7.87 (2H, s), 7.68 and 7.18 (4H, ABq, J=8.7Hz), 7.35 (1H, t, J=7.9Hz), 6.81 (1H, d, J=9.4Hz), 6.72 (1H, s), 6.71 (1H, d, J=6.8Hz), 4.80 (2H, s), 4.29 (1H, brt, J=12.2Hz), 4.10 (1H, t, J=6.7Hz), 2.43 (1H, t, J=6.7Hz), 2.39-2.19 (2H, m), 1.97-1.78 (4H, m), 1.76 (3H, s), 1.70-1.56 (1H, m), 1.43-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	497 (M+1)	

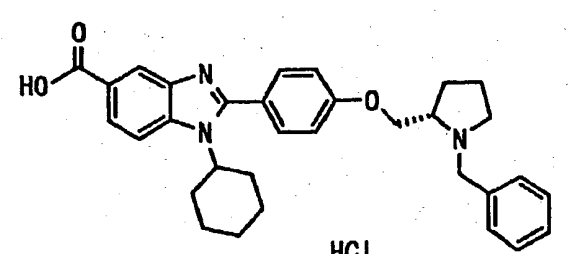
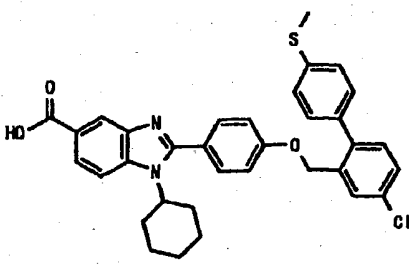
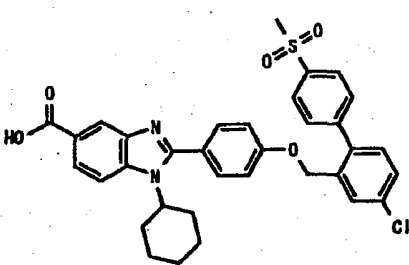
Example No.	165	1H NMR (δ) ppm
 HCl		300MHz, DMSO-d6 11.21 (1H, brs), 8.33 (1H, s), 8.25 (1H, d, J=8.6Hz), 8.04 (1H, d, J=8.6Hz), 7.78 (2H, d, J=8.7Hz), 7.70-7.67 (2H, m), 7.55-7.42 (3H, m), 7.27 (2H, d, J=8.7Hz), 4.73-4.30 (5H, m), 4.20-3.97 (1H, m), 3.42-3.10 (2H, m), 2.45-1.23 (14H, m)
Purity	> 90% (NMR)	
MS		

Table 46

Example No.	166	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 8.13 (1H, d, J=8.4Hz), 7.97 (1H, d, J=9.0Hz), 7.73 (1H, d, J=1.8Hz), 7.68 (2H, d, J=8.4Hz), 7.54 (1H, d, J=8.4, 2.1Hz), 7.41-7.31 (5H, m), 7.19 (2H, d, J=8.4Hz), 5.10 (2H, s), 4.32 (1H, m), 2.50 (3H, s), 2.40-2.15 (2H, m), 2.10-1.75 (4H, m), 1.75-1.55 (1H, m), 1.55-1.10 (3H, m).
Purity	> 90% (NMR)	
MS	583 (M+1)	

Example No.	167	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.25 (1H, s), 8.09 (1H, d, J=8.4Hz), 8.00 (2H, d, J=8.4Hz), 7.94 (1H, d, J=8.7Hz), 7.80 (1H, d, J=2.1Hz), 7.73 (2H, d, J=8.1Hz), 7.65 (2H, d, J=8.7Hz), 7.60 (1H, dd, J=8.1, 2.1Hz), 7.44 (1H, d, J=8.1Hz), 7.16 (2H, d, J=8.7Hz), 5.13 (2H, s), 4.30 (1H, m), 3.26 (3H, s), 2.40-1.15 (2H, m), 2.05-1.75 (4H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	615 (M+1)	

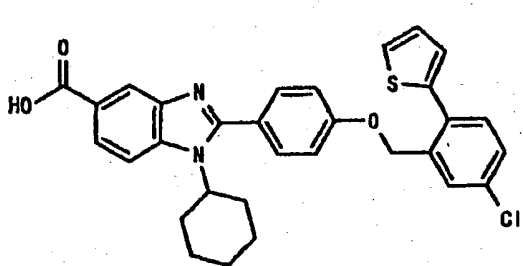
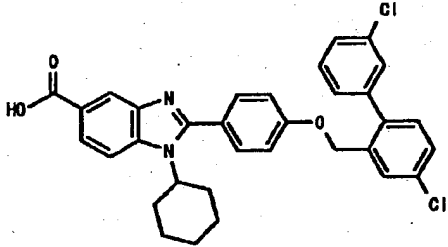
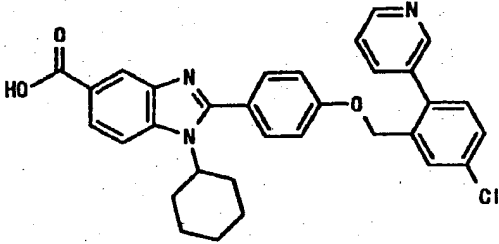
Example No.	168	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.1 (1H, brs), 8.32 (1H, s), 8.28 (1H, d, J=8.8Hz), 8.05 (1H, d, J=8.7Hz), 7.80-7.75 (3H, m), 7.69 (1H, d, J=4.1Hz), 7.57 (2H, m), 7.34-7.29 (3H, m), 7.20-7.15 (1H, m), 5.24 (2H, s), 4.39 (1H, m), 2.45-2.20 (2H, m), 2.20-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	543 (M+1)	

Table 47

Example No.	169	1H NMR( $\delta$ ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31(1H, s), 8.26(1H, d, J=8.7Hz), 8.05(1H, d, J=8.7Hz), 7.78-7.71(3H, m), 7.59-7.41(6H, m), 7.23(2H, d, J=9.0Hz), 5.11(2H, s), 4.35(1H, m), 2.40-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-1.15(3H, m).
Purity	> 90% (NMR)	
MS	571(M+1)	

Example No.	170	1H NMR( $\delta$ ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.7(1H, brs), 8.66(1H, s), 8.61(1H, m), 8.21(1H, s), 7.92-7.79(4H, m), 7.61-7.56(3H, m), 7.50-7.43(2H, m), 7.10(2H, d, J=8.7Hz), 5.09(2H, s), 4.26(1H, m), 2.40-2.15(2H, m), 2.00-1.75(4H, m), 1.75-1.55(1H, m), 1.50-1.15(3H, m).
Purity	> 90% (NMR)	
MS	538(M+1)	

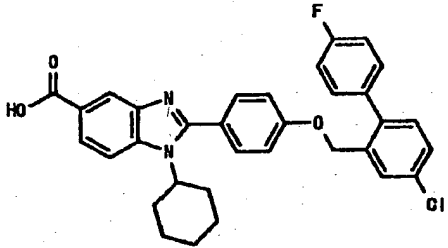
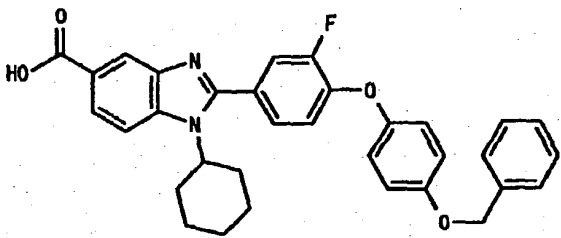
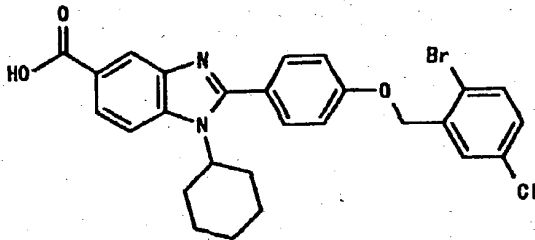
Example No.	171	1H NMR( $\delta$ ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31(1H, s), 8.25(1H, d, J=8.7Hz), 8.04(1H, d, J=8.7Hz), 7.74-7.71(3H, m), 7.57-7.46(3H, m), 7.39(1H, d, J=8.1Hz), 7.31-7.21(4H, m), 5.11(2H, s), 4.35(1H, m), 2.40-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-1.15(3H, m).
Purity	> 90% (NMR)	
MS	555(M+1)	

Table 48

Example No.	172	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, s), 7.99 (1H, d, J=8.7Hz), 7.88 (1H, d, J=10.5Hz), 7.70 (1H, dd, J=11.4, 1.8Hz), 7.48-7.32 (6H, m), 7.17-7.09 (5H, m), 5.12 (2H, s), 4.30 (1H, m), 2.40-2.15 (2H, m), 2.05-1.75 (4H, m), 1.75-1.55 (1H, m), 1.55-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	537 (M+1)	

Example No.	173	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.29 (1H, d, J=8.7Hz), 8.06 (1H, d, J=8.7Hz), 7.82-7.74 (4H, m), 7.45 (1H, dd, J=8.4, 3.0Hz), 7.39 (2H, d, J=8.7Hz), 5.28 (2H, s), 4.40 (1H, m), 2.40-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	540 (M+1)	

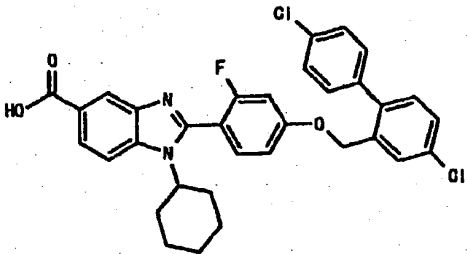
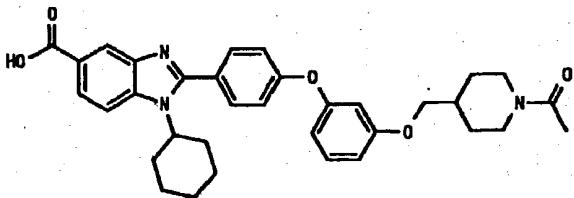
Example No.	174	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.80 (1H, brs), 8.26 (1H, s), 8.01 (1H, d, J=8.7Hz), 7.85 (1H, d, J=8.7Hz), 7.80-7.70 (1H, m), 7.60-7.36 (7H, m), 7.18-6.91 (2H, m), 5.09 (2H, s), 4.11-3.90 (1H, m), 2.32-1.18 (14H, m)
Purity	> 90% (NMR)	
MS	590 (M+1)	

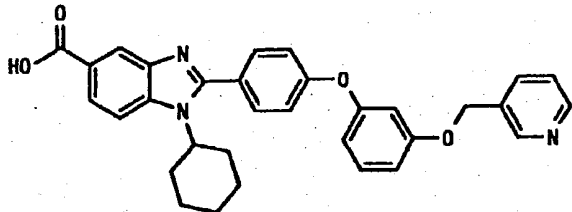
Table 49

Example No.	175	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.75 (1H, s), 8.21 (1H, s), 7.94 and 7.85 (2H, ABq, J=8.7 Hz), 7.61 and 7.00 (4H, A'B'q, J=8.5 Hz), 7.31-6.91 (2H, m), 7.25 (2H, d, J=7.7 Hz), 5.41 (2H, brs), 4.54 (2H, d, J=6.6 Hz), 4.35-4.14 (2H, m), 2.49-2.15 (3H, m), 1.95-1.55 (5H, m), 1.50-1.13 (5H, m), 1.10-0.77 (2H, m)
Purity	> 90% (NMR)	
MS	568 (M+1)	

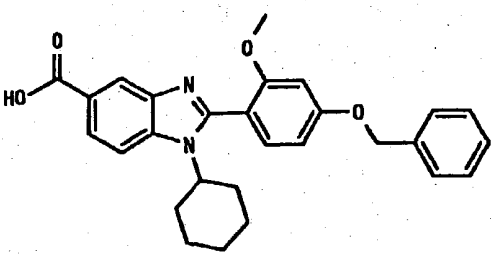
Example No.	176	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.24 (1H, s), 7.97 and 7.87 (2H, ABq, J=8.6Hz), 7.69 and 7.19 (4H, A' B' q, J=8.6Hz), 7.35 (1H, t, J=8.1Hz), 6.81 (1H, d, J=9.2Hz), 6.72 (1H, s), 6.71 (1H, d, J=6.5Hz), 4.48-4.20 (2H, m), 3.95-3.75 (3H, m), 3.03 (1H, t, J=12.3Hz), 2.60-2.40 (1H, m), 2.39-2.15 (2H, m), 2.07-1.58 (6H, m), 1.99 (3H, s), 1.50-1.00 (5H, m)
Purity	> 90% (NMR)	
MS	568 (M+1)	

Example No.	177	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.76 (1H, s), 8.23 (1H, s), 7.96 and 7.86 (2H, ABq, J=8.6 Hz), 7.69 and 7.20 (4H, A'B'q, J=8.6 Hz), 7.39 (1H, t, J=8.2 Hz), 6.86 (1H, d, J=8.3 Hz), 6.81 (1H, s), 6.76 (1H, d, J=8.0 Hz), 4.83 (2H, s), 4.31 (1H, brt, J=12.2 Hz), 2.39-2.19 (2H, m), 1.99-1.79 (4H, m), 1.70-1.58 (1H, m), 1.48-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	467 (M+1)	

Table 50

Example No.	178	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.85 (1H, s), 8.75 (1H, s), 8.63 (2H, d, J=3.8Hz), 8.25 (1H, s), 8.04-8.01 (2H, m), 8.02 and 7.90 (2H, ABq, J=8.6Hz), 7.72 and 7.20 (4H, A'B'q, J=8.6Hz), 7.57 (2H, dd, J=7.8, 5.0Hz), 7.40 (1H, t, J=8.2Hz), 6.93 (1H, d, J=8.2Hz), 6.87 (1H, s), 6.77 (1H, d, J=8.2Hz), 5.23 (2H, s), 4.33 (1H, br t, J=12.2Hz), 2.40-2.18 (2H, m), 2.00-1.55 (5H, m), 1.50-1.15 (2H, m)
Purity	> 90% (NMR)	
MS	520 (M+1)	

Example No.	179	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.29 (1H, d, J=9.0Hz), 8.06 (1H, d, J=8.7Hz), 7.61 (1H, d, J=8.4Hz), 7.58-7.32 (5H, m), 6.98 (1H, d, J=2.1Hz), 6.93 (1H, dd, J=8.7, 2.1Hz), 5.27 (2H, s), 4.16-4.00 (1H, m), 3.87 (3H, s), 2.20-2.12 (2H, m), 2.02-1.98 (4H, m), 1.70-1.60 (1H, m), 1.52-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	457 (M+1)	

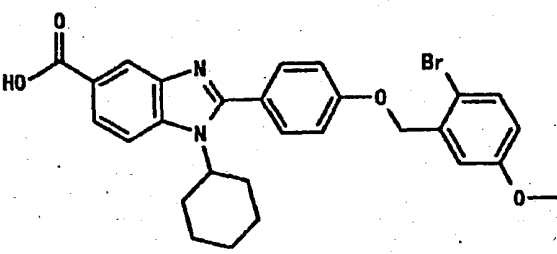
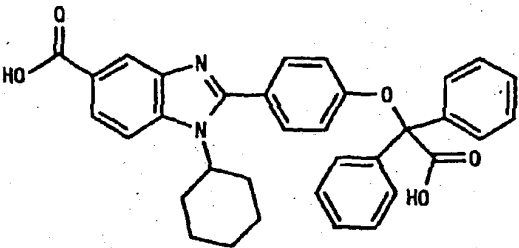
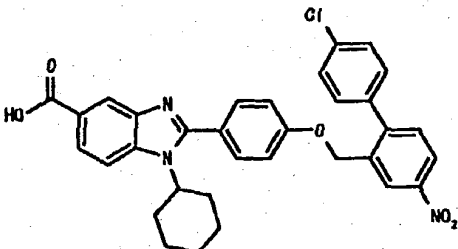
Example No.	180	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 7.91 (1H, d, J=8.6Hz), 7.85 (1H, d, J=8.6Hz), 7.63 (2H, d, J=8.4Hz), 7.60 (1H, d, J=9.0Hz), 7.25 (2H, d, J=8.4Hz), 7.23 (1H, d, J=3.0Hz), 6.95 (1H, dd, J=9.0, 3.0Hz), 5.19 (2H, s), 4.30 (1H, m), 3.78 (3H, s), 2.40-2.19 (2H, m), 2.00-1.87 (4H, m), 1.66 (1H, m), 1.49-1.18 (3H, m)
Purity	> 90% (NMR)	
MS	536 (M+1)	

Table 51

Example No.	181	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.19 (1H, s), 7.95 (1H, d, J=8.7Hz), 7.86 (1H, d, J=8.7Hz), 7.65 (4H, d, J=7.4Hz), 7.47 (2H, d, J=8.7Hz), 7.44-7.27 (6H, m), 6.99 (2H, d, J=8.7Hz), 4.20 (1H, m), 2.34-2.12 (2H, m), 1.98-1.75 (4H, m), 1.64 (1H, m), 1.46-1.13 (3H, m).
Purity	> 90% (NMR)	
MS	547 (M+)	

Example No.	182	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.55 (1H, d, J=2.1Hz), 8.32 (1H, m), 8.21 (1H, s), 7.95 (1H, d, J=8.4Hz), 7.86 (1H, d, J=7.8Hz), 7.68-7.56 (7H, m), 7.14 (2H, d, J=8.7Hz), 5.21 (1H, s), 4.26 (1H, m), 2.35-2.15 (2H, m), 2.00-1.75 (4H, m), 1.74-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	582 (M+)	

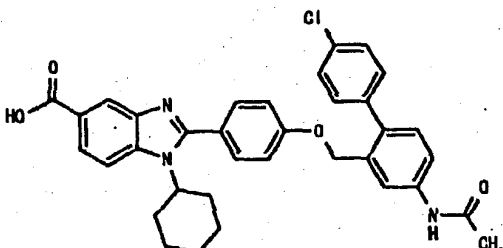
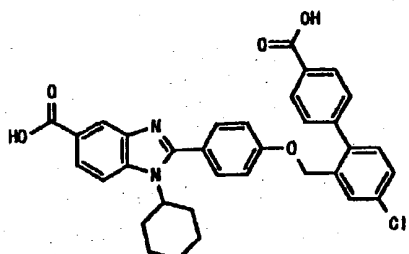
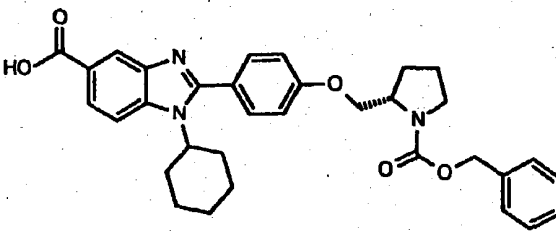
Example No.	183	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.16 (1H, s), 8.25 (1H, s), 8.07 (1H, d, J=8.7Hz), 7.94-7.87 (2H, m), 7.71-7.62 (3H, m), 7.50-7.42 (4H, m), 7.30 (1H, d, J=8.4Hz), 7.14 (2H, d, J=8.4Hz), 5.06 (2H, s), 4.31 (1H, m), 2.35-2.15 (2H, m), 2.05-1.75 (4H, m), 1.75-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	594 (M+)	

Table 52

Example No.	184	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.2 (2H, brs), 8.30 (1H, s), 8.26 (1H, d, J=8.8Hz), 8.04 (1H, d, J=8.8Hz), 8.00 (2H, d, J=8.2Hz), 7.79 (1H, s), 7.73 (2H, d, J=8.7Hz), 7.61-7.56 (3H, m), 7.44 (1H, d, J=8.3Hz), 7.23 (2H, d, J=8.8Hz), 5.13 (2H, s), 4.35 (1H, m), 2.45-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (1H, m), 1.75-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	581 (M+1)	

Example No.	185	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, m), 8.24 (1H, d, J=9.0Hz), 8.03 (1H, d, J=9.0Hz), 7.79-7.10 (9H, m), 5.20-5.07 (2H, m), 4.43-4.04 (4H, m), 3.50-3.36 (2H, m), 2.40-1.19 (14H, m)
Purity	> 90% (NMR)	
MS	554 (M+1)	

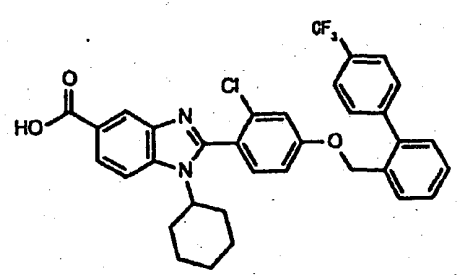
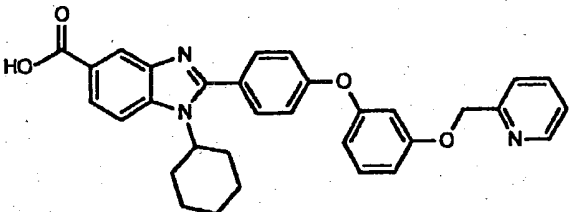
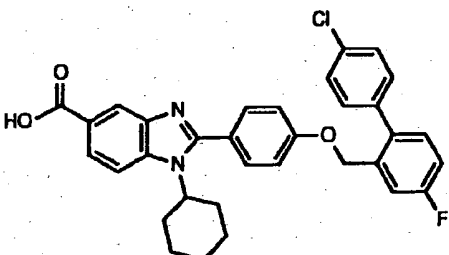
Example No.	186	1H NMR (δ) ppm
		(DMSO-d6) δ : 8.29 (1H, brs), 8.10 (1H, d, J=8.4Hz), 7.97 (1H, d, J=8.4Hz), 7.79 (2H, d, J=8.4Hz), 7.74-7.67 (1H, m), 7.68 (2H, d, J=8.4Hz), 7.61 (1H, d, J=8.4Hz), 7.57-7.50 (2H, m), 7.46-7.39 (1H, m), 7.29 (1H, d, J=2.4Hz), 7.11 (1H, dd, J=2.4, 8.4Hz), 5.12 (2H, s), 3.99-3.84 (1H, m), 2.35-1.72 (6H, m), 1.68-1.55 (1H, m), 1.42-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	605 (M+1)	



Table 53

Example No.	187	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.76 (1H, s), 8.57 (1H, d, J=4.4Hz), 8.23 (1H, s), 7.96 and 7.86 (2H, ABq, J=8.2Hz), 7.87-7.82 (1H, m), 7.68 and 7.12 (4H, A'B'q, J=8.6Hz), 7.53 (2H, d, J=7.8Hz), 7.37 (1H, t, J=8.3Hz), 7.36-7.33 (1H, m), 6.90 (1H, d, J=8.3Hz), 6.83 (1H, s), 6.74 (1H, d, J=8.0Hz), 5.20 (2H, s), 4.31 (1H, br t, J=12.2Hz), 2.35-2.19 (2H, m), 1.99-1.57 (5H, m), 1.45-1.20 (2H, m)
Purity	> 90% (NMR)	
MS	520 (M+1)	

Example No.	188	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.77 (1H, brs), 8.21 (1H, d, J=1.4Hz), 7.92 (1H, d, J=8.7Hz), 7.88 (1H, dd, J=8.7, 1.4Hz), 7.57 (2H, d, J=8.7Hz), 7.57-7.27 (7H, m), 7.11 (2H, d, J=8.7Hz), 5.07 (2H, s), 4.26 (1H, m), 2.36-2.16 (2H, m), 1.98-1.75 (4H, m), 1.64 (1H, m), 1.49-1.17 (3H, m).
Purity	> 90% (NMR)	
MS	555 (M+1)	

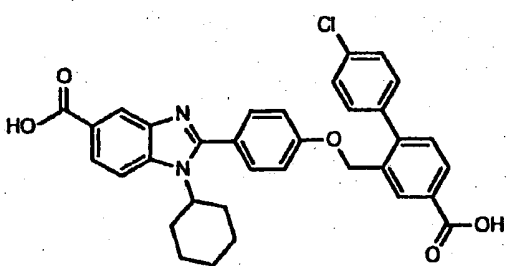
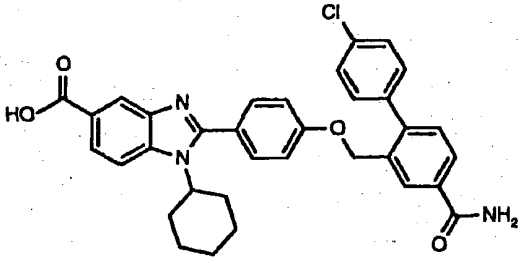
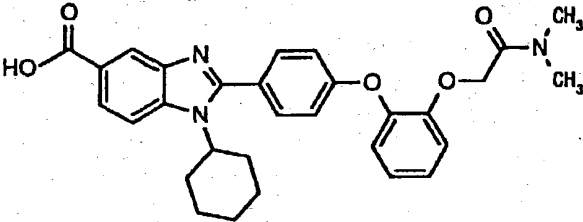
Example No.	189	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.30-8.20 (2H, m), 8.10-7.98 (2H, m), 7.74 (2H, d, J=9.0Hz), 7.60-7.46 (5H, m), 7.24 (2H, d, J=9.0Hz), 5.19 (2H, s), 4.44-4.30 (1H, m), 2.40-2.20 (2H, m), 2.12-1.78 (4H, m), 1.72-1.58 (4H, m)
Purity	> 90% (NMR)	
MS	581 (M+1)	

Table 54

Example No.	190	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.36-7.90 (5H, m), 7.74 (2H, d, J=8.6Hz), 7.60-7.40 (5H, m), 7.25 (2H, d, J=8.7Hz), 5.14 (2H, s), 4.45-4.28 (1H, m), 2.40-2.15 (4H, m), 1.75-1.55 (1H, m), 1.55-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	580 (M+1)	

Example No.	191	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.94 (1H, d, J=8.4Hz), 7.85 (1H, d, J=8.7Hz), 7.61 (2H, d, J=8.7Hz), 7.25-7.00 (6H, m), 4.86 (2H, s), 4.30 (1H, m), 2.89 (3H, s), 2.80 (3H, s), 2.29 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	514 (M+1)	

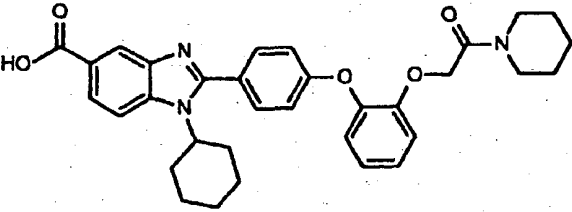
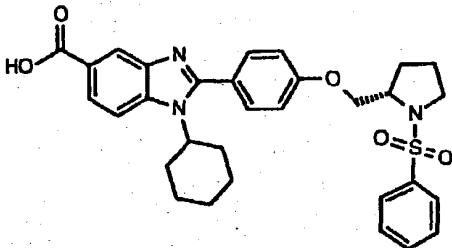
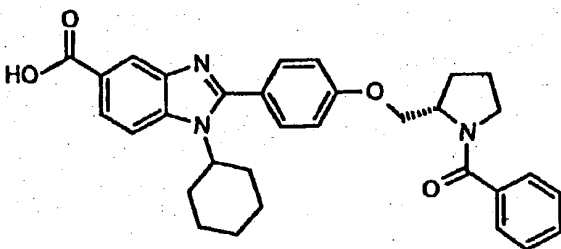
Example No.	192	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.94 (1H, d, J=8.4Hz), 7.85 (1H, d, J=8.7Hz), 7.61 (2H, d, J=8.7Hz), 7.26-7.01 (6H, m), 4.84 (2H, s), 4.31 (1H, m), 3.36 (4H, m), 2.29 (2H, m), 2.00-1.75 (4H, m), 1.75-1.15 (10H, m)
Purity	> 90% (NMR)	
MS	554 (M+1)	

Table 55

Example No.	193	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 13.00 (1H, brs), 8.29 (1H, d, J=1.4Hz), 8.15 (1H, d, J=8.8Hz), 7.97 (1H, dd, J=1.4Hz, 8.8Hz), 7.89 (2H, d, J=8.8Hz), 7.80-7.60 (5H, m), 7.25 (2H, d, J=8.8Hz), 4.47-3.90 (4H, m), 3.20-3.10 (2H, m), 2.41-1.22 (14H, m)
Purity	> 90% (NMR)	
MS	560 (M+1)	

Example No.	194	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.80 (1H, brs), 8.23 (1H, s), 7.97 (1H, d, J=8.5Hz), 7.87 (1H, d, J=8.5Hz), 7.70-7.17 (9H, m), 4.60-4.13 (4H, m), 3.72-3.40 (2H, m), 2.40-1.15 (14H, m)
Purity	> 90% (NMR)	
MS	524 (M+1)	

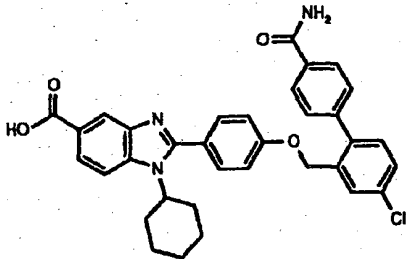
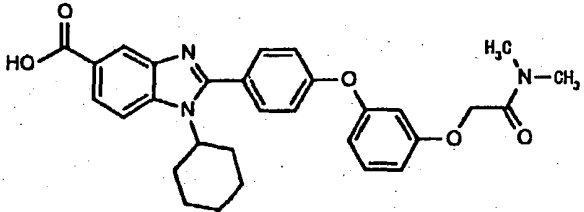
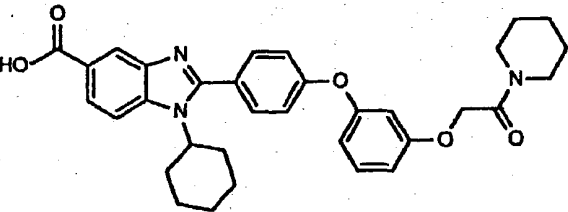
Example No.	195	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.25 (1H, s), 8.09-7.92 (5H, m), 7.77 (1H, s), 7.65 (2H, d, J=8.4Hz), 7.59-7.51 (3H, m), 7.43 (2H, d, J=8.4Hz), 7.17 (2H, d, J=8.7Hz), 5.10 (2H, s), 4.30 (1H, m), 2.40-2.15 (2H, m), 2.10-1.75 (4H, m), 1.75-1.55 (1H, m), 1.55-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	580 (M+1)	

Table 56

Example No.	196	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.95 (1H, d, J=8.4Hz), 7.86 (1H, d, J=8.4Hz), 7.69 and 7.18 (4H, ABq, J=8.7Hz), 7.34 (1H, t, J=8.0Hz), 6.80-6.69 (3H, m), 4.83 (2H, s), 4.31 (1H, m), 2.98 (3H, s), 2.84 (3H, s), 2.29 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	514 (M+1)	

Example No.	197	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.95 (1H, d, J=8.4Hz), 7.86 (1H, d, J=8.7Hz), 7.69 and 7.18 (4H, ABq, J=8.7Hz), 7.35 (1H, t, J=8.4Hz), 6.80-6.70 (3H, m), 4.82 (2H, s), 4.31 (1H, m), 3.40 (4H, m), 2.29 (2H, m), 2.00-1.75 (4H, m), 1.70-1.15 (10H, m)
Purity	> 90% (NMR)	
MS	554 (M+1)	

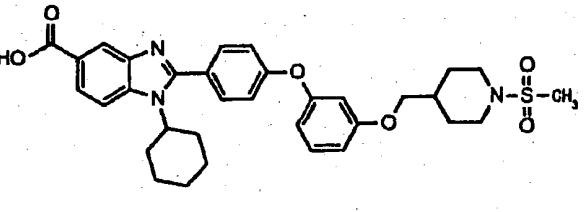
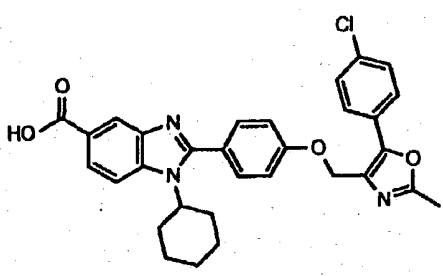
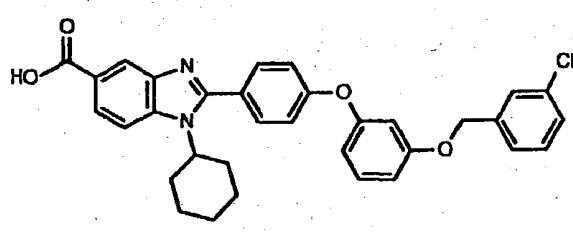
Example No.	198	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, s), 8.23 (1H, d, J=4.4Hz), 7.95 and 7.86 (2H, ABq, J=8.6Hz), 7.69 and 7.19 (4H, A'B'q, J=8.6Hz), 7.36 (1H, t, J=7.8Hz), 6.82 (1H, d, J=9.3Hz), 6.73 (1H, s), 6.71 (1H, d, J=7.2Hz), 4.30 (1H, brt, J=12.2Hz), 3.89 (2H, d, J=6.0Hz), 3.59 (2H, d, J=11.7Hz), 2.85 (3H, s), 2.73 (2H, t, J=10.5Hz), 2.41-2.20 (2H, m), 1.98-1.59 (8H, m), 1.46-1.12 (5H, m)
Purity	> 90% (NMR)	
MS	604 (M+1)	

Table 57

Example No.	199	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.30 (1H, d, J=8.9Hz), 8.06 (1H, d, J=8.7Hz), 7.79 (2H, d, J=8.7Hz), 7.70 (2H, d, J=8.7Hz), 7.61 (2H, d, J=8.7Hz), 7.39 (2H, d, J=8.8Hz), 5.28 (2H, s), 4.39 (1H, m), 2.50-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	542 (M+1)	

Example No.	200	1H NMR (δ) ppm
		(DMSO-d6) δ : 8.23 (1H, s), 7.96 (1H, d, J=8.6Hz), 7.86 (1H, d, J=8.6Hz), 7.69 (2H, d, J=8.4Hz), 7.52 (1H, s), 7.50-7.30 (4H, m), 7.18 (2H, d, J=8.4Hz), 6.90 (1H, d, J=8.3Hz), 6.84 (1H, s), 6.74 (1H, d, J=8.3Hz), 5.15 (2H, s), 4.39-4.21 (1H, m), 2.39-2.18 (2H, m), 1.99-1.80 (4H, m), 1.71-1.59 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

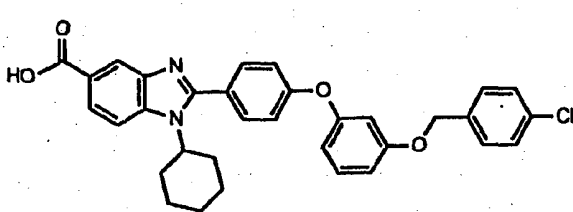
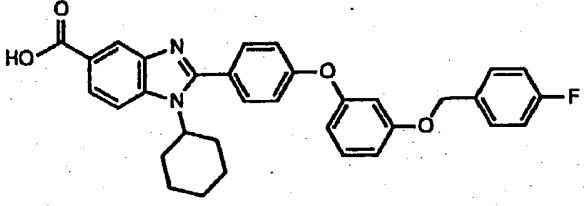
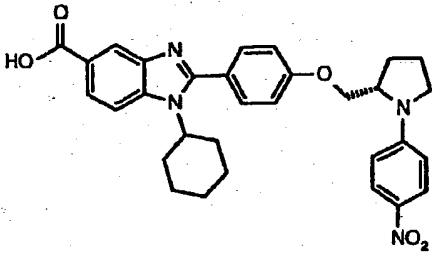
Example No.	201	1H NMR (δ) ppm
		(DMSO-d6) δ : 8.26 (1H, s), 8.06 (1H, d, J=8.7Hz), 7.92 (1H, d, J=8.7Hz), 7.72 (2H, d, J=8.7Hz), 7.47 (4H, s), 7.38 (1H, t, J=8.2Hz), 7.20 (2H, d, J=8.7Hz), 6.90 (1H, d, J=8.2Hz), 6.83 (1H, s), 6.74 (1H, d, J=8.2Hz), 5.14 (2H, s), 2.40-2.19 (2H, m), 2.04-1.78 (4H, m), 1.71-1.60 (1H, m), 1.50-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

Table 58

Example No.	202	1H NMR(δ) ppm
		(DMSO-d6) δ : 12.81 (1H, brs), 8.24 (1H, s), 7.99 (1H, d, J=8.7Hz), 7.87 (1H, d, J=8.7Hz), 7.69 (2H, d, J=8.6Hz), 7.53-7.47 (2H, m), 7.38 (1H, t, J=8.2Hz), 7.26-7.16 (4H, m), 6.89 (1H, d, J=8.2Hz), 6.82 (1H, s), 6.73 (1H, d, J=8.2Hz), 5.11 (2H, s), 4.40-4.21 (1H, m), 2.40-2.17 (2H, m), 2.01-1.77 (4H, m), 1.71-1.59 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	537 (M+1)	

Example No.	203	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.74 (1H, brs), 8.21 (1H, s), 8.08 (2H, d, J=9.0Hz), 7.93 (1H, d, J=8.7Hz), 7.85 (2H, d, J=8.7Hz), 7.58 (2H, d, J=8.7Hz), 7.13 (2H, d, J=8.7Hz), 6.83 (2H, d, J=9.0Hz), 4.50-4.08 (4H, m), 3.68-3.30 (2H, m), 2.40-1.23 (14H, m)
Purity	> 90% (NMR)	
MS	541 (M+1)	

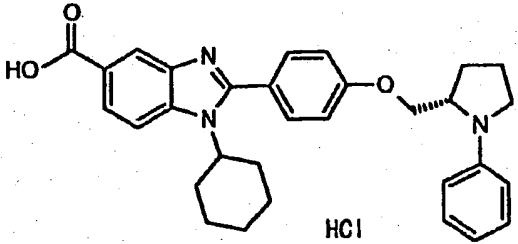
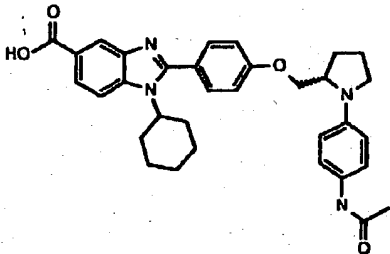
Example No.	204	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.39-8.28 (2H, m), 8.08 (1H, d, J=8.8Hz), 7.76 (2H, d, J=8.7Hz), 7.29 (2H, d, J=8.7Hz), 7.25-7.13 (2H, m), 6.80-6.60 (3H, m), 4.46-3.98 (4H, m), 3.51-3.42 (1H, m), 3.20-3.04 (1H, m), 2.39-1.20 (14H, m)
Purity	> 90% (NMR)	
MS		

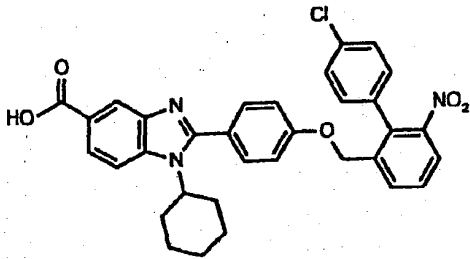
Table 59

Example No.	205	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9. 59 (1H, brs), 8. 23 (1H, s), 8. 04 (1H, d, J=8. 4Hz), 7. 90 ( 1H, d, J=8. 4Hz), 7. 62 (2H, d, J=8. 7Hz), 7. 39 (2H, 2H, d, J= 8. 7Hz) 7. 18 (2H, d, J=8. 7Hz) , 6. 63 (2H, d, J=8. 7Hz), 3. 95 -3. 37 (4H, m), 3. 51-3. 40 (1H , m), 3. 17-3. 02 (1H. m), 2. 39 -1. 18 (17H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

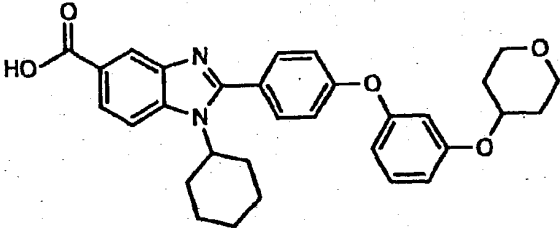
Example No.	206	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 13.1 (1H, brs), 8.33 (1H, s), 8.29 (1H, d, J=8.8Hz), 8.06 (1H, d, J=8.7Hz), 7.77 (2H, d, J=8.7Hz), 7.59-7.52 (4H, m), 7.35 (2H, d, J=8.8Hz), 5.19 (2H, s), 4.39 (1H, m), 2.71 (3H, s), 2.45-2.20 (2H, m), 2.20-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	558 (M+1)	

Example No.	207	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.29 (1H, s), 8.26 (1H, d, J=8.8Hz), 8.04 (1H, d, J=8.7Hz), 7.73 (2H, d, J=8.8Hz), 7.50-7.41 (6H, m), 7.36 (2H, d, J=8.8Hz), 7.18-7.13 (2H, m), 6.84 (1H, s), 4.33 (1H, m), 2.40-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	539 (M+1)	

Table 60

Example No.	208	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.32(1H, s), 8.27(1H, d, J=9.0Hz), 8.07-8.00(3H, m), 7.79-7.70(3H, m), 7.51(2H, d, J=8.1Hz), 7.40(2H, d, J=8.4Hz), 7.18(2H, d, J=8.7Hz), 4.99(2H, s), 4.34(1H, m), 2.40-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-1.15(3H, m).
Purity	> 90% (NMR)	
MS	582 (M+1)	

Example No.	209	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.24(1H, d, J=4.4Hz), 7.98 and 7.88(2H, ABq, J=8.6Hz), 7.70 and 7.19(4H, A'B'q, J=8.4Hz), 7.35(1H, t, J=8.4Hz), 6.86(1H, d, J=8.1Hz), 6.79(1H, s), 6.71(1H, d, J=8.1Hz), 4.65-4.53(1H, m), 4.31(1H, brt, J=12.2Hz), 3.88-3.78(2H, m), 3.48(2H, t, J=9.0Hz), 2.39-2.19(2H, m), 1.02-1.71(6H, m), 1.70-1.50(3H, m), 1.46-1.19(3H, m)
Purity	> 90% (NMR)	
MS	513 (M+1)	

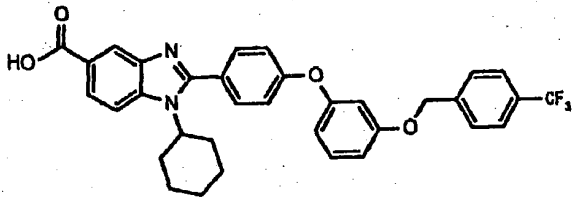
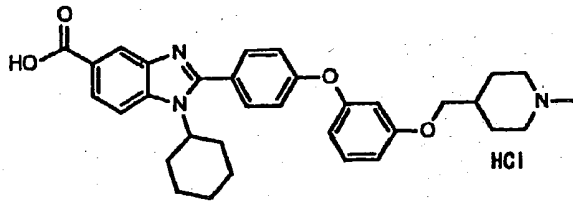
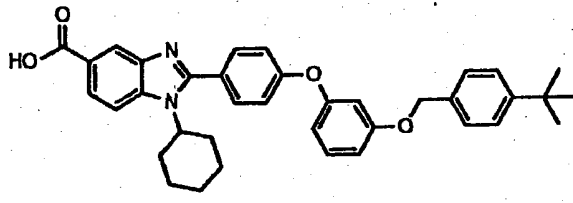
Example No.	210	1H NMR(δ) ppm
		300MHz, DMSO-d6 12.75(1H, s), 8.23(1H, s), 7.96 and 7.87(2H, ABq, J=8.7Hz), 7.84-7.66(6H, m), 7.38(1H, t, J=8.4Hz), 7.18(2H, d, J=8.4Hz), 6.91(1H, d, J=9.0Hz), 6.84(1H, s), 6.74(1H, d, J=8.1Hz), 5.26(2H, s), 4.31(1H, brt, J=12.2Hz), 2.40-2.20(2H, m), 1.99-1.76(4H, m), 1.69-1.58(1H, m), 1.45-1.20(3H, m)
Purity	> 90% (NMR)	
MS	587 (M+1)	



Table 61

Example No.	211	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.15 and 7.47 (2H, ABq, J=9.0Hz), 7.77 and 7.24 (4H, ABq, J=8.9Hz), 7.39 (1H, t, J=7.8Hz), 6.84 (1H, d, J=9.3Hz), 6.76 (1H, s), 6.75 (1H, d, J=9.5Hz), 4.36 (1H, brt, J=12.2Hz), 3.89 (2H, d, J=6.0Hz), 3.42 (2H, d, J=10.8Hz), 3.04-2.88 (2H, m), 2.78-2.60 (1H, m), 2.71 (2H, d, J=4.8Hz), 2.38-2.20 (2H, m), 2.07-1.80 (7H, m), 1.70-1.20 (5H, m)
Purity	> 90% (NMR)	
MS	540 (M+1)	

Example No.	212	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.93 and 7.87 (2H, ABq, J=8.6Hz), 7.68 and 7.17 (4H, A'B'q, J=8.7Hz), 7.43-7.33 (5H, m), 6.87 (1H, d, J=8.1Hz), 7.18 (2H, d, J=8.4Hz), 6.91 (1H, d, J=9.0Hz), 6.81 (1H, s), 6.72 (1H, d, J=8.0Hz), 5.08 (2H, s), 4.36 (1H, brt, J=12.2Hz), 2.37-2.20 (2H, m), 1.98-1.78 (4H, m), 1.69-1.60 (1H, m), 1.41-1.21 (3H, m), 1.28 (9H, s)
Purity	> 90% (NMR)	
MS	575 (M+1)	

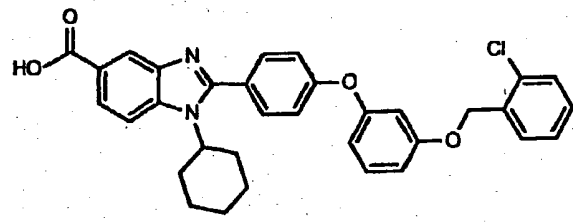
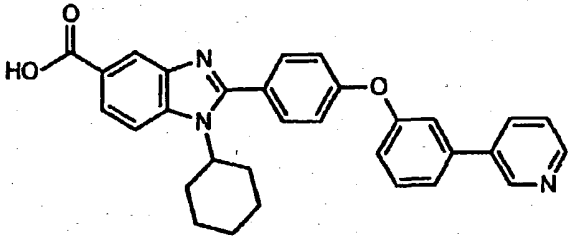
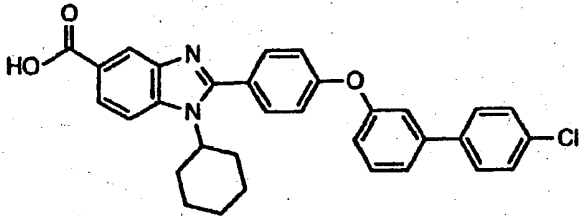
Example No.	213	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.95 and 7.86 (2H, ABq, J=8.4Hz), 7.69 and 7.19 (4H, A'B'q, J=8.7Hz), 7.62-7.36 (5H, m), 6.90 (1H, d, J=8.1Hz), 6.84 (1H, s), 6.76 (1H, d, J=8.1Hz), 5.19 (2H, s), 4.31 (1H, brt, J=12.2Hz), 2.40-2.19 (2H, m), 1.99-1.76 (4H, m), 1.68-1.55 (1H, m), 1.50-1.18 (3H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

Table 62

Example No.	214	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.94 (1H, d, J=2.1Hz), 8.60 (1H, dd, J=4.8, 1.5Hz), 8.23 (1H, d, J=1.5Hz), 8.12 (1H, dt, J=8.1, 2.1Hz), 7.93 (1H, d, J=8.7Hz), 7.87 (1H, dd, J=8.7, 1.5Hz), 7.70 (1H, d, J=8.7Hz), 7.67-7.54 (3H, m), 7.50 (1H, dd, J=8.1, 4.8Hz), 7.25 (2H, d, J=8.7Hz), 7.21 (1H, m), 4.31 (1H, m), 2.38-2.19 (2H, m), 2.00-1.78 (4H, m), 1.65 (1H, m), 1.48-1.22 (3H, m).
Purity	> 90% (NMR)	
MS	490 (M+1)	

Example No.	215	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.75 (1H, brs), 8.23 (1H, s), 7.95 (1H, d, J=8.7Hz), 7.86 (1H, d, J=8.7Hz), 7.73 (2H, d, J=8.4Hz), 7.71 (2H, d, J=8.4Hz), 7.63-7.39 (2H, m), 7.52 (2H, d, J=8.4Hz), 7.24 (2H, d, J=8.4Hz), 7.18 (1H, m), 4.31 (1H, m), 2.39-2.20 (2H, m), 2.00-1.76 (4H, m), 1.65 (1H, m), 1.49-1.18 (3H, m).
Purity	> 90% (NMR)	
MS	523 (M+1)	

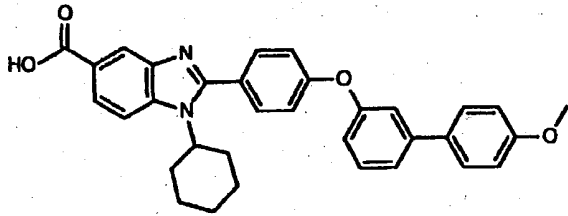
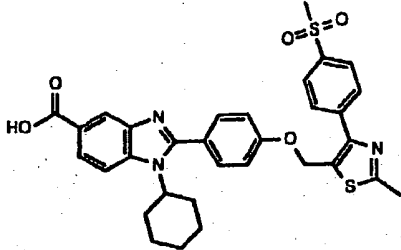
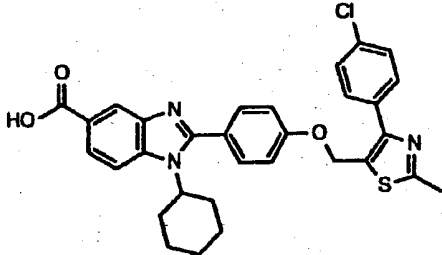
Example No.	216	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.77 (1H, s), 8.23 (1H, d, J=1.4Hz), 7.95 (1H, d, J=8.6Hz), 7.86 (1H, dd, J=8.6, 1.4Hz), 7.70 (2H, d, J=8.7Hz), 7.64 (2H, d, J=8.8Hz), 7.56-7.48 (2H, m), 7.40 (1H, s), 7.23 (2H, d, J=8.7Hz), 7.10 (1H, m), 7.03 (2H, d, J=8.8Hz), 4.31 (1H, m), 3.80 (3H, s), 2.48-2.20 (2H, m), 2.00-1.88 (4H, m), 1.66 (1H, m), 1.50-1.21 (3H, m).
Purity	> 90% (NMR)	
MS	519 (M+1)	

Table 63

Example No.	217	1H NMR ( $\delta$ ) ppm
		(DMSO-d <sub>6</sub> ) $\delta$ : 12.80 (1H, brs), 8.23 (1H, s), 8.04 (1H, d, J=8.6 Hz), 7.96 (3H, d, J=8.6 Hz), 7.86 (1H, d, J=8.7 Hz), 7.63 (2H, d, J=8.6 Hz), 7.25 (2H, d, J=8.6 Hz), 5.50 (2H, s), 4.36-4.21 (1H, m), 3.27 (3H, s), 2.74 (3H, s), 2.40-2.19 (2H, m), 1.99-1.79 (4H, m), 1.71-1.60 (1H, m), 1.49-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	602 (M+1)	

Example No.	218	1H NMR ( $\delta$ ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.9 (1H, brs), 8.25 (1H, s), 8.04 (1H, d, J=8.7 Hz), 7.91 (1H, d, J=8.6 Hz), 7.72 (2H, d, J=8.5 Hz), 7.67 (2H, d, J=8.7 Hz), 7.56 (2H, d, J=8.5 Hz), 7.26 (2H, d, J=8.7 Hz), 5.45 (2H, s), 4.31 (1H, m), 2.71 (3H, s), 2.40-2.15 (2H, m), 2.05-1.80 (4H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	558 (M+1)	

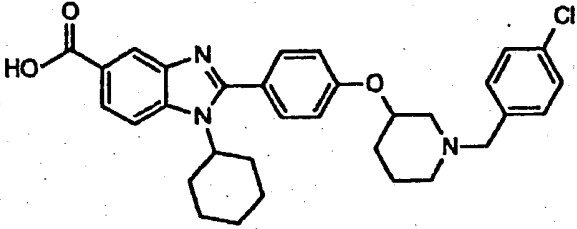
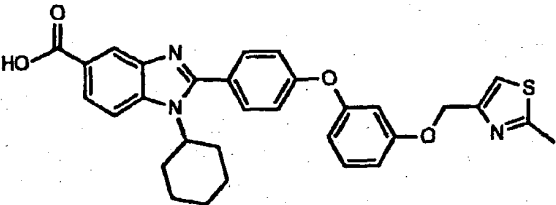
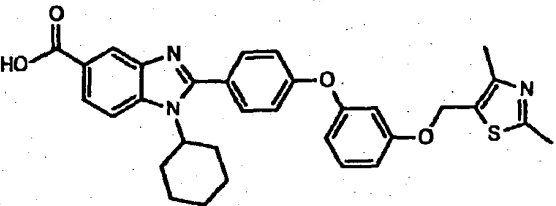
Example No.	219	1H NMR ( $\delta$ ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.21 (1H, d, J=1.5 Hz), 7.93 (1H, d, J=9.0 Hz), 7.84 (1H, dd, J=9.0, 1.5 Hz), 7.56 (2H, d, J=8.7 Hz), 7.42-7.30 (4H, m), 7.12 (2H, d, J=8.7 Hz), 4.53 (1H, brs), 4.36-4.20 (1H, m), 3.55 (2H, brs), 3.00-2.90 (1H, m), 2.70-2.58 (1H, m), 2.40-1.10 (18H, m)
Purity	> 90% (NMR)	
MS	544 (M+1)	

Table 64

Example No.	220	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.76 (1H, s), 8.23 (1H, s), 7.96 and 7.87 (2H, ABq, J=8.9 Hz), 7.69 and 7.19 (4H, A'B'q, J=8.6 Hz), 7.55 (1H, s), 7.37 (1H, t, J=8.1 Hz), 6.91 (1H, d, J=7.8 Hz), 6.85 (1H, s), 6.74 (1H, d, J=7.5 Hz), 5.13 (2H, s), 4.31 (1H, brt, J=12.2 Hz), 2.65 (3H, s), 2.41-2.20 (2H, m), 2.00-1.74 (4H, m), 1.70-1.59 (1H, m), 1.58-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	540 (M+1)	

Example No.	221	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.96 and 7.86 (2H, ABq, J=8.6 Hz), 7.69 and 7.18 (4H, A'B'q, J=8.7 Hz), 7.37 (1H, t, J=8.2 Hz), 6.87 (1H, d, J=8.2 Hz), 6.82 (1H, s), 6.75 (1H, d, J=8.0 Hz), 5.24 (2H, s), 4.32 (1H, brt, J=12.2 Hz), 2.58 (3H, s), 2.38-2.20 (2H, m), 2.30 (3H, s), 2.00-1.79 (4H, m), 1.70-1.59 (1H, m), 1.44-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	554 (M+1)	

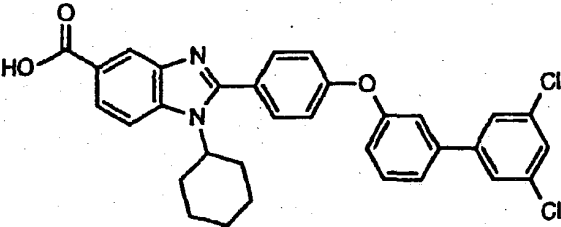
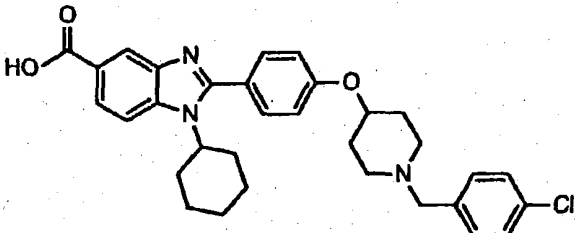
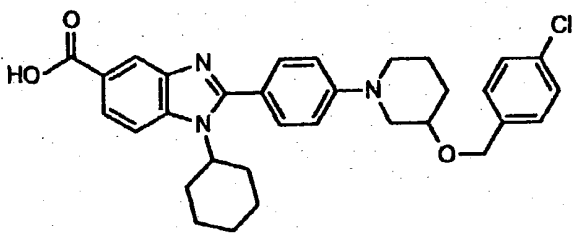
Example No.	222	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.88 (1H, brs), 8.25 (s, 1H), 8.07-7.57 (11H, m), 7.26 (2H, d, J=8.7 Hz), 7.24 (1H, m), 4.34 (1H, m), 2.30-2.20 (2H, m), 2.03-1.78 (4H, m), 1.64 (1H, m), 1.49-1.19 (3H, m).
Purity	> 90% (NMR)	
MS	557 (M+1)	

Table 65

Example No.	223	1H NMR(δ) ppm
		300MHz, DMSO-d6 10.96(1H, brs), 8.21(1H, d, J=1.4Hz), 7.93(1H, d, J=8.7 Hz), 7.84(1H, dd, J=8.7, 1.4 Hz), 7.76-7.40(7H, m), 7.18(2H, d, J=8.0Hz), 4.24-4.16(2H, m), 2.40-1.12(18H, m)
Purity	>90% (NMR)	
MS	544 (M+1)	

Example No.	224	1H NMR(δ) ppm
		(DMSO-d6) δ : 8.22(1H, s), 8.07(1H, d, J=8.4Hz), 7.92(1H, d, J=8.4Hz), 7.54(2H, d, J=8.7Hz), 7.40(2H, d, J=8.4Hz), 7.30(2H, d, J=8.4Hz), 7.14(2H, d, J=8.7Hz), 4.61(2H, s), 4.48-4.32(1H, m), 3.82(1H, brd, J=12.3Hz), 3.65-3.47(2H, m), 3.10(brdd, J=8.4, 12.3Hz), 2.40-2.20(2H, m), 2.09-1.76(6H, m), 1.71-1.16(6H, m)
Purity	>90% (NMR)	
MS	544 (M+1)	

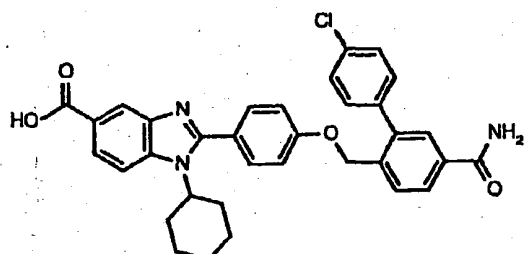
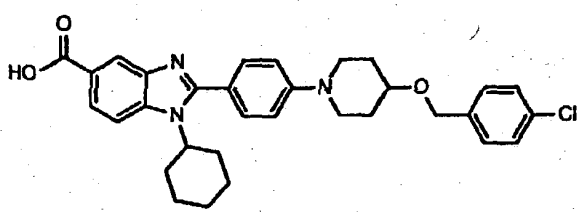
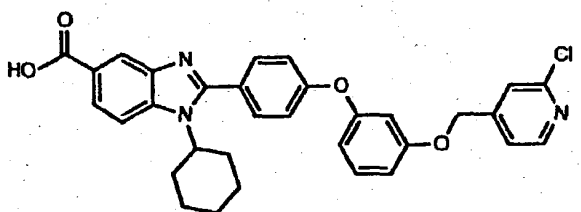
Example No.	225	1H NMR(δ) ppm
		(DMSO-d6) δ : 12.83(1H, brs), 8.21(1H, s), 8.10(1H, brs), 7.01-7.91(2H, m), 7.89-7.82(2H, m), 7.75(1H, d, J=8.0Hz), 7.59(2H, d, J=8.7Hz), 7.53(4H, s), 7.46(1H, brs), 7.12(2H, d, J=8.7Hz), 7.23(2H, s), 4.35-4.17(1H, m), 2.38-2.20(2H, m), 1.99-1.79(4H, m), 1.71-1.59(1H, m), 1.48-1.18(3H, m)
Purity	>90% (NMR)	
MS	580 (M+1)	

Table 66

Example No.	226	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 and 8.08 (2H, ABq, J=8.7 Hz), 8.31 (1H, m), 7.66 and 7.26 (4H, A'B'q, J=9.2 Hz), 7.42 and 7.39 (4H, A''B''q, J=8.7 Hz), 4.57 (2H, s), 4.50 (1H, br t, J=12.2 Hz), 3.85-3.62 (3H, m), 3.28-3.16 (2H, m), 2.42-2.23 (2H, m), 2.14-1.81 (6H, m), 1.72-1.25 (6H, m)
Purity	> 90% (NMR)	
MS	544 (M+1)	

Example No.	227	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.43 (1H, d, J=5.0 Hz), 8.23 (1H, s), 7.96 and 7.86 (2H, ABq, J=8.6 Hz), 7.69 and 7.18 (4H, A'B'q, J=8.6 Hz), 7.57 (1H, s), 7.47 (1H, d, J=5.0 Hz), 7.40 (2H, t, J=8.2 Hz), 6.91 (1H, d, J=8.3 Hz), 6.85 (1H, s), 6.77 (1H, d, J=7.9 Hz), 5.25 (2H, s), 4.31 (1H, br t, J=12.2 Hz), 2.40-2.19 (2H, m), 1.99-1.75 (4H, m), 1.73-1.57 (1H, m), 1.49-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	554 (M+1)	

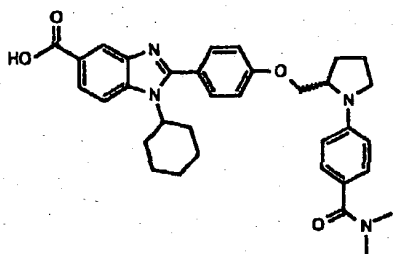
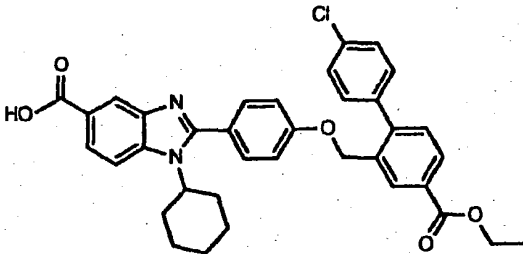
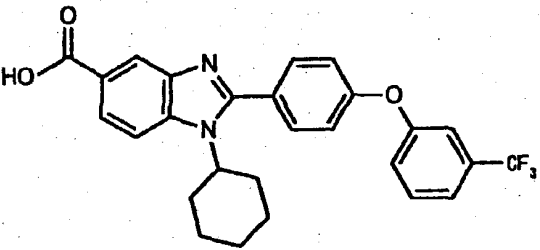
Example No.	228	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.80 (1H, brs), 8.22 (1H, s), 7.94 (1H, d, J=8.6 Hz), 7.87 (1H, d, J=8.6 Hz), 7.60 (2H, d, J=8.7 Hz), 7.32 (2H, d, J=8.7 Hz), 7.17 (2H, d, J=8.7 Hz), 6.70 (2H, d, J=8.7 Hz), 4.35-3.97 (4H, m), 3.62-3.11 (2H, m), 2.96 (6H, s), 2.39-1.12 (14H, m)
Purity	> 90% (NMR)	
MS	567 (M+1)	

Table 67

Example No.	229	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.25(1H, s), 8.20(1H, s), 8.04(1H, dd, J=8.1, 1.8Hz), 7.92(1H, d, J=8.1Hz), 7.84(1H, d, J=9.9Hz), 7.62-7.50(7H, m), 7.12(2H, d, J=8.7Hz), 5.14(2H, s), 4.36(2H, q, J=6.9Hz), 4.30-4.20(1H, m), 2.38-2.18(2H, m), 1.98-1.18(8H, m), 1.35(3H, t, J=6.9Hz)
Purity	>90% (NMR)	
MS	608 (M+1)	

Example No.	230	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.35(1H, s), 8.27(1H, d, J=8.7Hz), 8.05(1H, d, J=9.0Hz), 7.87(2H, d, J=8.7Hz), 7.74(1H, t, J=8.1Hz), 7.64(1H, d, J=7.8Hz), 7.59-7.50(2H, m), 7.36(2H, d, J=8.7Hz), 4.39(1H, m), 2.40-2.15(2H, m), 2.15-1.95(2H, m), 1.95-1.75(2H, m), 1.75-1.55(1H, m), 1.55-1.20(3H, m).
Purity	about 90% (NMR)	
MS	481 (M+1)	

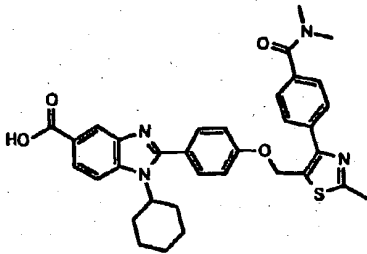
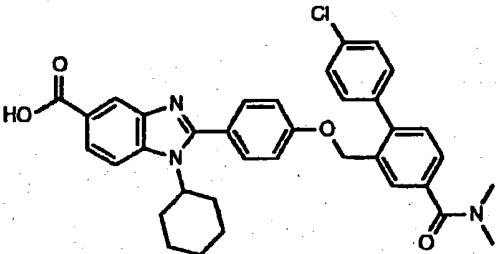
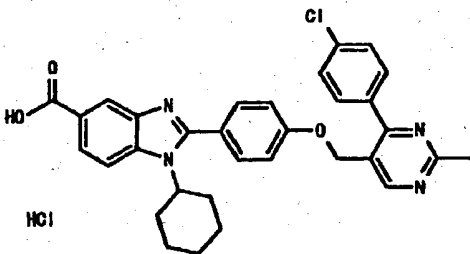
Example No.	231	1H NMR(δ) ppm
		300MHz DMSO-d6 12.78(1H, brs), 8.23(1H, d, J=1.5Hz), 7.96(1H, d, J=8.7Hz), 7.87(1H, dd, J=8.7, 1.5Hz), 7.75(2H, d, J=8.4Hz), 7.63(2H, d, J=8.4Hz), 7.52(2H, d, J=8.4Hz), 7.24(2H, d, J=8.4Hz), 5.47(2H, s), 4.29(1H, m), 2.97(6H, brs), 2.72(3H, s), 2.39-2.16(2H, m), 2.00-1.78(4H, m), 1.71-1.59(1H, m), 1.49-1.17(3H, m).
Purity	about 90% (NMR)	
MS	595 (M+1)	

Table 68

Example No.	232	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.8 (1H, brs), 8.22 (1H, s), 7.96 (1H, d, J=8.7Hz), 7.86 (1H, d, J=8.6Hz), 7.70 (1H, s), 7.59 (2H, d, J=8.7Hz), 7.53-7.50 (5H, m), 7.42 (1H, d, J=7.9Hz), 7.12 (2H, d, J=8.7Hz), 5.11 (2H, s), 4.27 (1H, m), 3.01 (3H, brs), 2.97 (3H, brs), 2.40-2.15 (2H, m), 2.00-1.75 (4H, m), 1.75-1.55 (1H, m), 1.50-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	608 (M+1)	

Example No.	233	<sup>1</sup> H NMR (δ) ppm
		DMSO-d <sub>6</sub> 13.20 (1H, brs), 8.99 (1H, s), 8.32 (1H, s), 8.25 (1H, d, J=8.8Hz), 8.04 (1H, d, J=8.6Hz), 7.79-7.74 (4H, m), 7.60 (2H, d, J=8.5Hz), 7.30 (2H, d, J=8.7Hz), 5.26 (2H, s), 4.36 (1H, m), 2.72 (3H, s), 2.50-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	553 (M+1-HCl)	

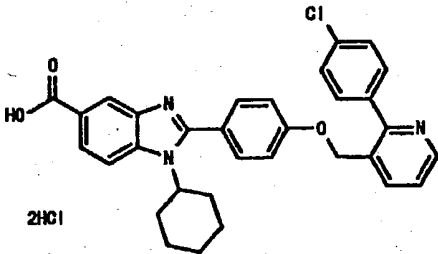
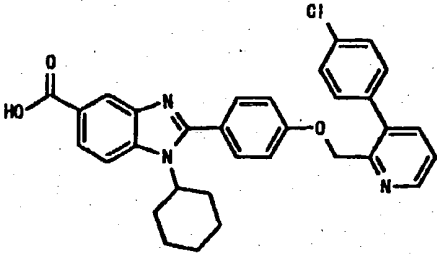
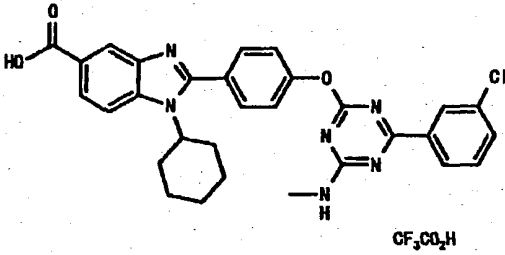
Example No.	234	<sup>1</sup> H NMR (δ) ppm
		DMSO-d <sub>6</sub> 8.77 (1H, d, J=3.6Hz), 8.36-8.26 (3H, m), 8.08 (1H, d, J=8.8Hz), 7.79 (2H, d, J=8.7Hz), 7.72-7.64 (3H, m), 7.58 (2H, d, J=8.4Hz), 7.30 (2H, d, J=8.7Hz), 5.26 (2H, s), 4.38 (1H, m), 2.50-2.15 (2H, m), 2.15-1.95 (2H, m), 1.95-1.75 (2H, m), 1.75-1.55 (1H, m), 1.55-1.15 (3H, m).
Purity	> 90% (NMR)	
MS	538 (M+1-2HCl)	



Table 69

Example No.	235	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.74 (1H, brs), 8.67 (1H, dd, J=3.1, 1.6Hz), 8.21 (1H, d, J=1.6Hz), 7.93 (1H, d, J=8.6Hz), 7.90-7.80 (2H, m), 7.60-7.50 (7H, m), 7.09 (2H, d, J=8.7Hz), 5.16 (2H, s), 4.26 (1H, m), 2.40-2.20 (2H, m), 2.00-1.60 (5H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	APCI-MS 538 (M+1)	

Example No.	236	1H NMR (δ) ppm
		300MHz, DMSO-d-6 8.40-7.40 (11H, m), 2.95, 2.81 (3H, each d, J=4.7Hz), 2.40-2.20 (2H, m), 2.10-1.80 (4H, m), 1.70-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	APCI-MS 555 (M+1)	

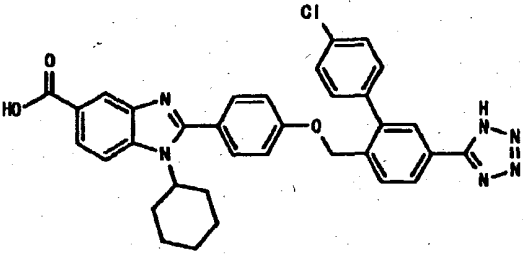
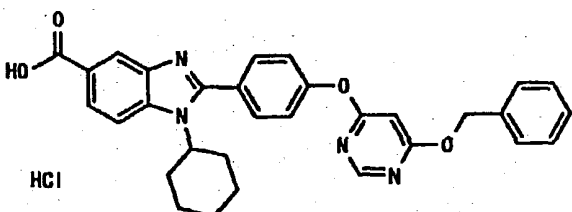
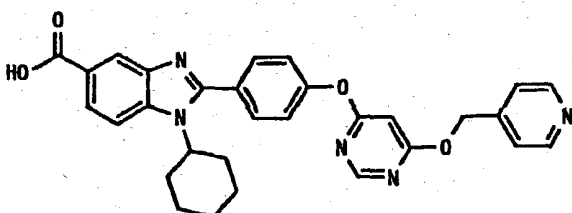
Example No.	237	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, s), 8.15 (1H, d, J=9.5Hz), 8.02 (1H, s), 8.00-7.80 (3H, m), 7.70-7.50 (6H, m), 7.12 (2H, d, J=8.7Hz), 5.16 (2H, s), 4.28 (1H, m), 2.40-2.20 (2H, m), 2.00-1.80 (4H, m), 1.65 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	FAB-MS 605 (M+1)	

Table 70

Example No.	238	1H NMR (δ) ppm
 HCl		300MHz, DMSO-d6 12.80 (1H, brs), 8.54 (1H, s), 8.25 (1H, s), 7.98 and 7.88 (2H, Abq, J=8.6Hz), 7.76 (2H, d, J=8.6Hz), 7.53-7.31 (3H, m), 6.61 (1H, s), 5.46 (2H, s), 4.32 (1H, brt), 2.40-2.20 (2H, m), 2.02-1.79 (4H, m), 1.69-1.59 (1H, m), 1.48-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	APCI-Ms 521 (M+1)	

Example No.	239	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.79 (1H, brs), 8.60 (2H, d, J=1.5Hz), 8.53 (1H, s), 8.25 (1H, s), 7.98 and 7.85 (2H, ABq, J=9.4Hz), 7.76 (2H, d, J=9.0Hz), 7.44 (4H, d, J=6.5Hz), 6.69 (1H, s), 5.53 (2H, s), 4.32 (1H, brt), 2.40-2.19 (2H, m), 2.03-1.82 (4H, m), 1.72-1.61 (1H, m), 1.42-1.22 (3H, m)
Purity	> 90% (NMR)	
MS	APCI-Ms 522 (M+1)	

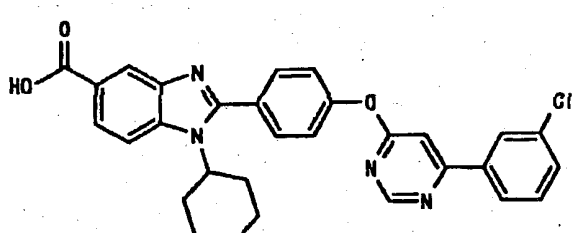
Example No.	240	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.90 (1H, s), 8.32 (1H, s), 8.28 (1H, s), 8.25 (1H, d, J=8.3Hz), 8.05 (1H, d, J=8.8Hz), 7.96 (1H, s), 7.93 (1H, d, J=8.8Hz), 7.83 (1H, d, J=8.4Hz), 7.68-7.59 (2H, m), 7.54 (2H, d, J=8.8Hz), 4.37 (1H, brt), 2.30 (2H, m), 2.00 (2H, m), 1.88 (2H, m), 1.67 (1H, m), 1.5-1.2 (3H, m)
Purity	> 90% (NMR)	
MS	APCI-Ms 525 (M+1)	

Table 71

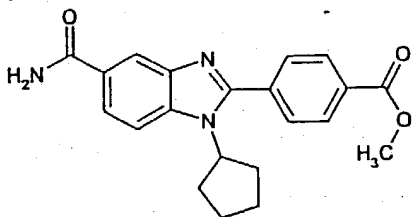
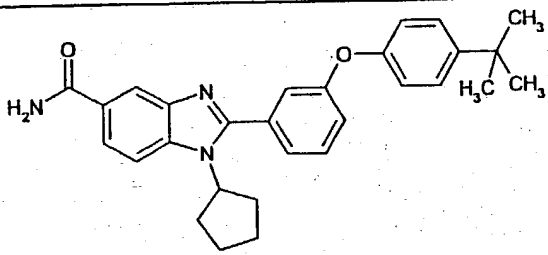
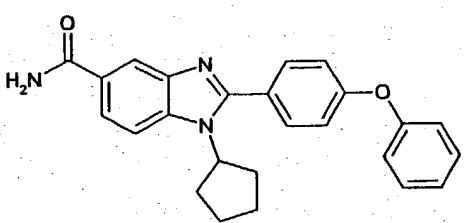
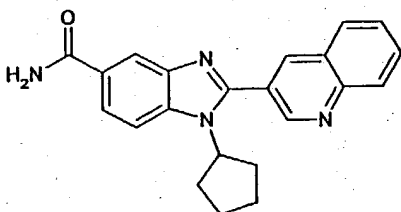
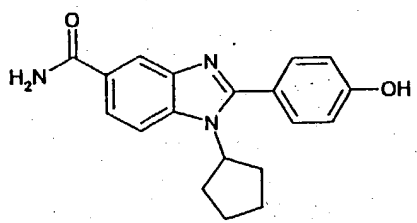
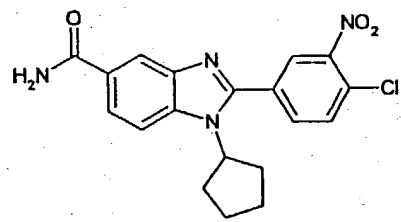
Ex. No.	Formula	MS
1001		364 (M+H)
1002		454 (M+H)
1003		398 (M+H)
1004		357 (M+H)
1005		322 (M+H)
1006		385 (M+H)

Table 72

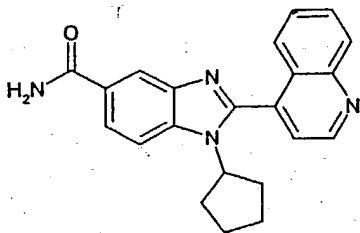
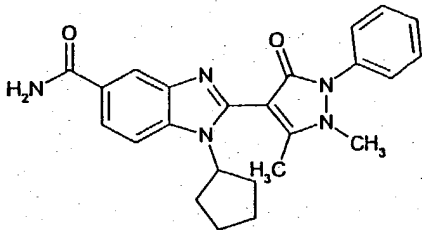
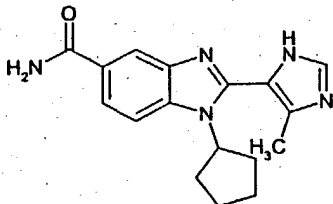
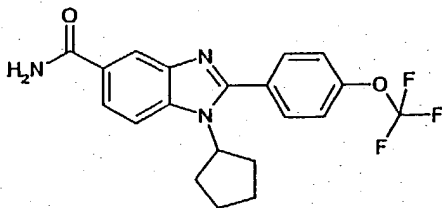
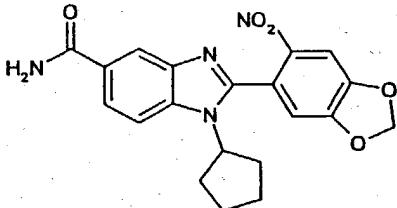
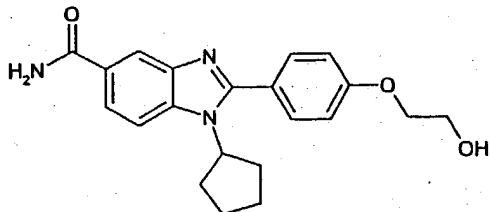
Ex. No.	Formula	MS
1007		357 (M+H)
1008		416 (M+H)
1009		310 (M+H)
1010		390 (M+H)
1011		395 (M+H)
1012		366 (M+H)

Table 73

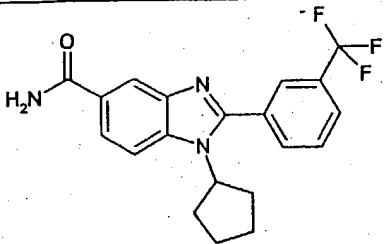
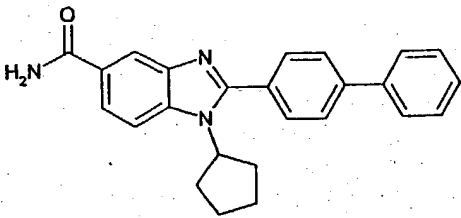
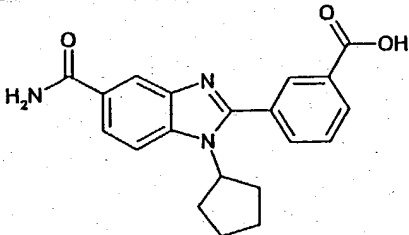
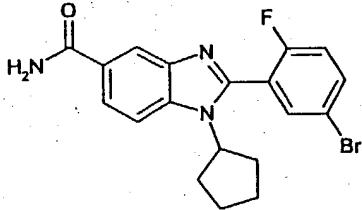
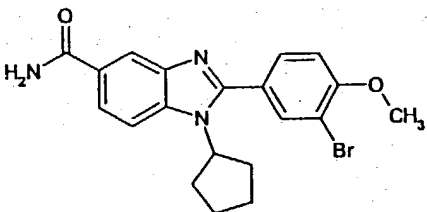
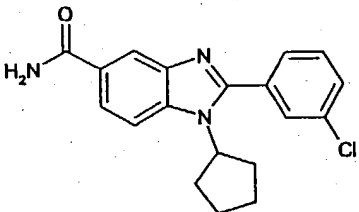
Ex. No.	Formula	MS
1013		374 (M+H)
1014		382 (M+H)
1015		350 (M+H)
1016		402 (M+H)
1017		414 (M+H)
1018		340 (M+H)

Table 74

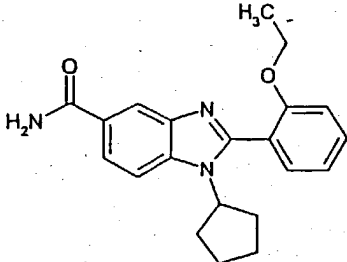
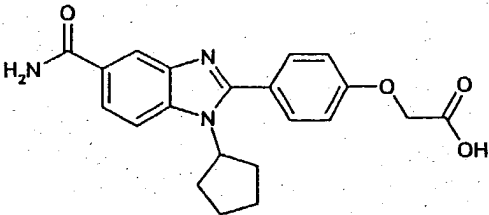
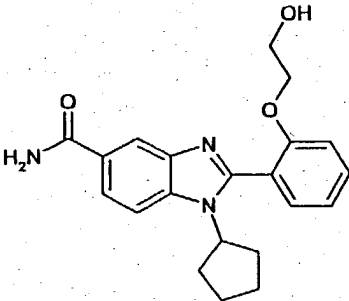
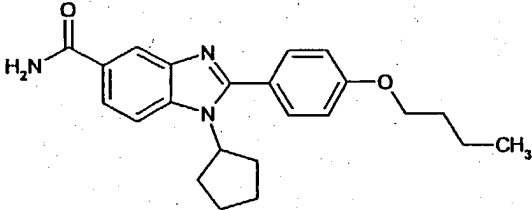
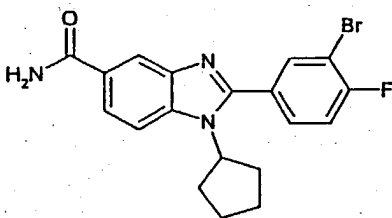
Ex. No.	Formula	MS
1019		350 (M+H)
1020		380 (M+H)
1021		366 (M+H)
1022		378 (M+H)
1023		402 (M+H)

Table 75

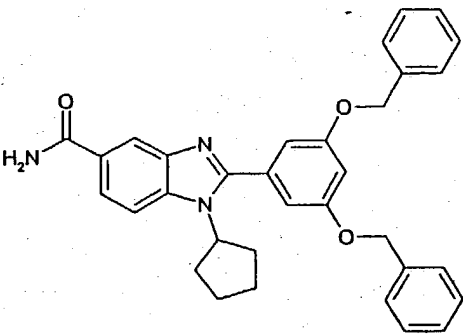
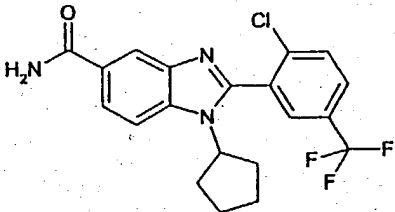
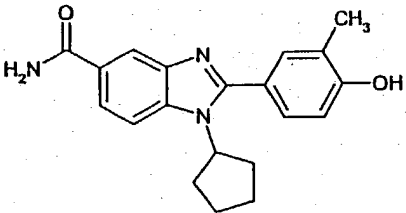
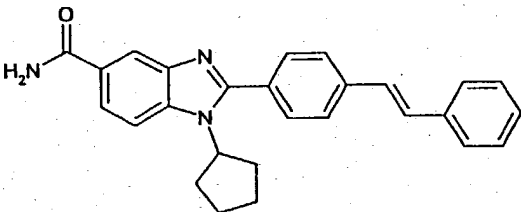
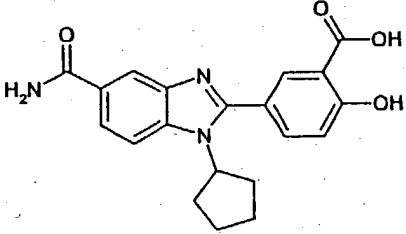
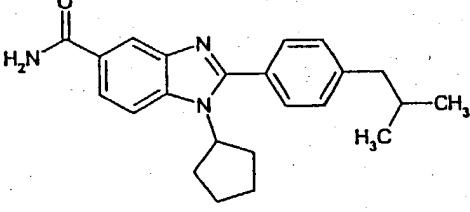
Ex. No.	Formula	MS
1024		518 (M+H)
1025		408 (M+H)
1026		336 (M+H)
1027		408 (M+H)
1028		366 (M+H)
1029		362 (M+H)

Table 76

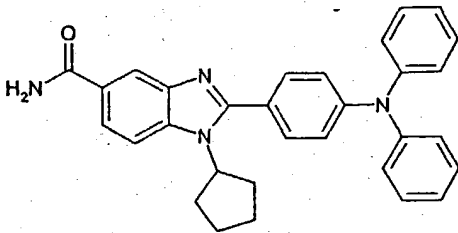
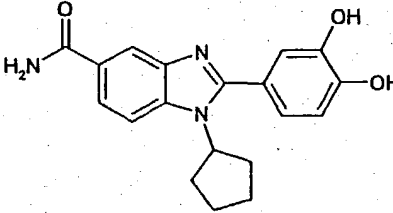
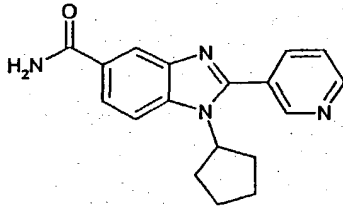
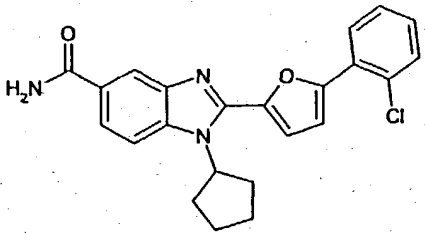
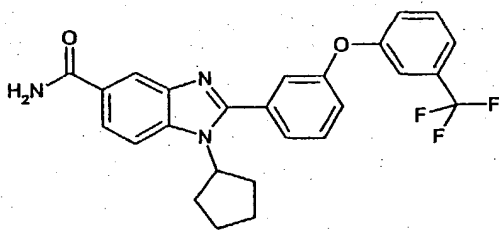
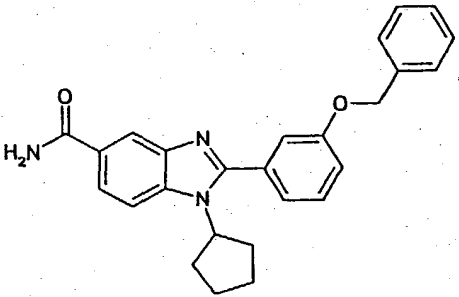
Ex. No.	Formula	MS
1030		473 (M+H)
1031		338 (M+H)
1032		307 (M+H)
1033		406 (M+H)
1034		466 (M+H)
1035		412 (M+H)



Table 77

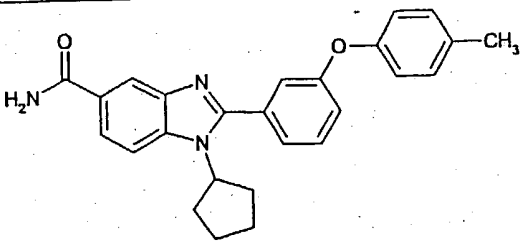
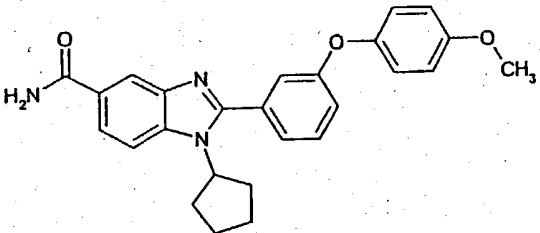
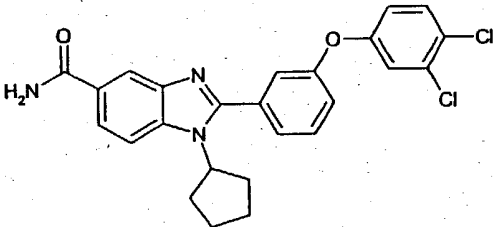
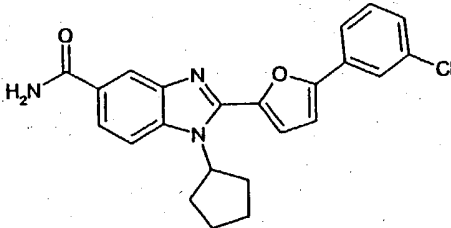
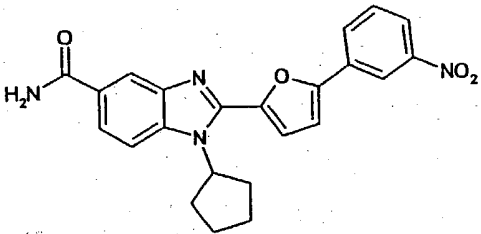
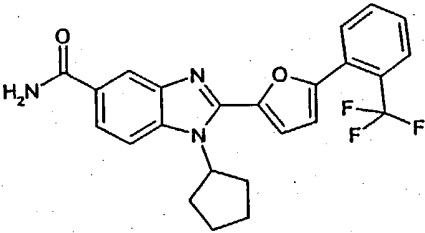
Ex. No.	Formula	MS
1036		412 (M+H)
1037		428 (M+H)
1038		466 (M+H)
1039		406 (M+H)
1040		417 (M+H)
1041		440 (M+H)

Table 78

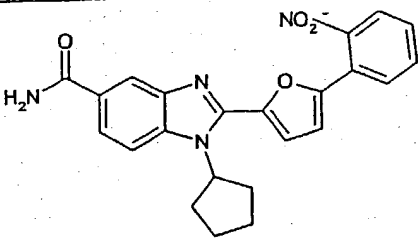
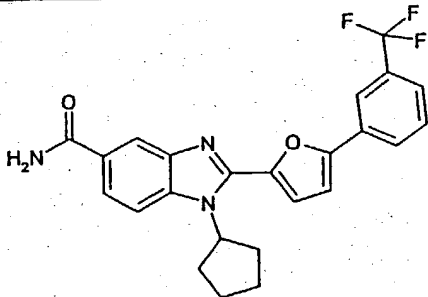
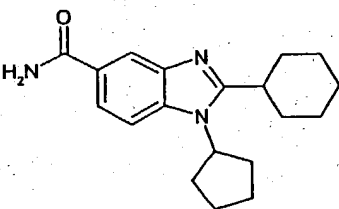
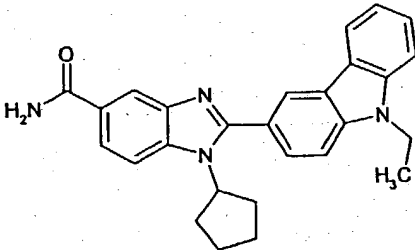
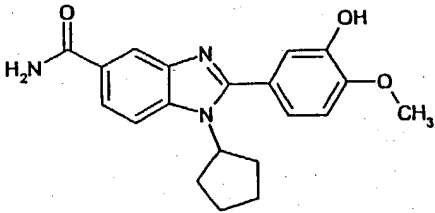
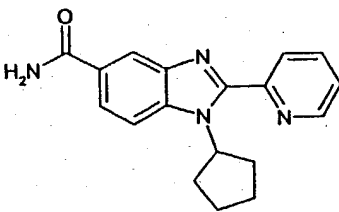
Ex. No.	Formula	MS
1042		417 (M+H)
1043		440 (M+H)
1044		312 (M+H)
1045		423 (M+H)
1046		352 (M+H)
1047		307 (M+H)

Table 79

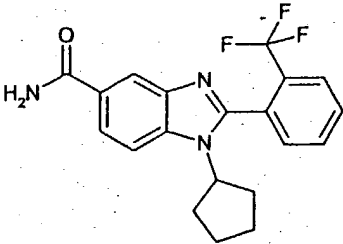
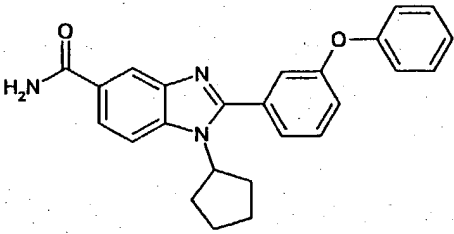
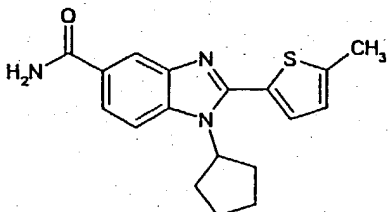
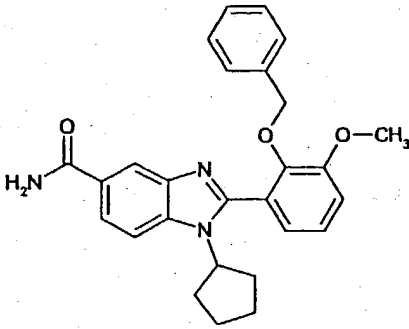
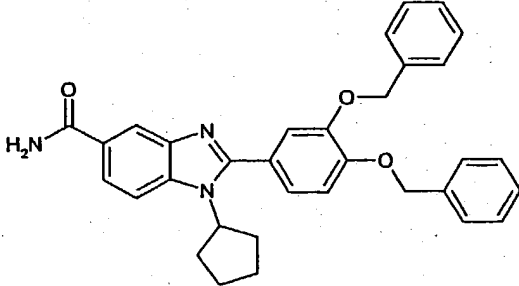
Ex. No.	Formula	MS
1048		374 (M+H)
1049		398 (M+H)
1050		326 (M+H)
1051		442 (M+H)
1052		518 (M+H)

Table 80

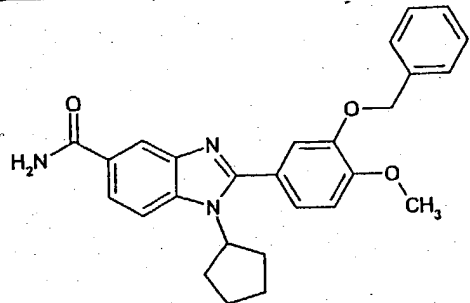
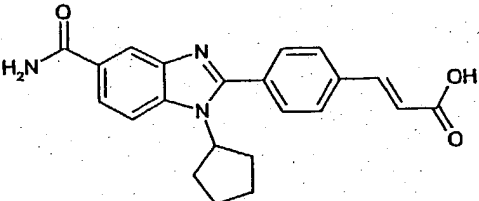
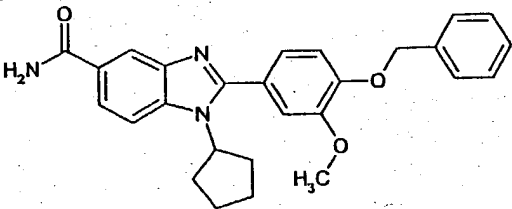
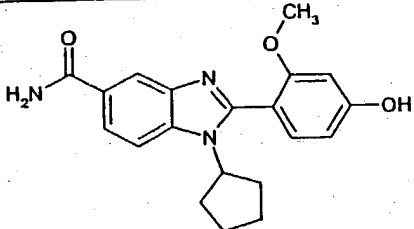
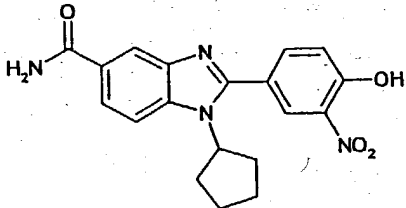
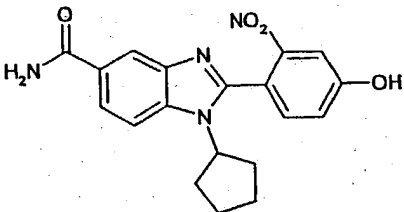
Ex. No.	Formula	MS
1053		442 (M+H)
1054		376 (M+H)
1055		442 (M+H)
1056		352 (M+H)
1057		367 (M+H)
1058		367 (M+H)

Table 81

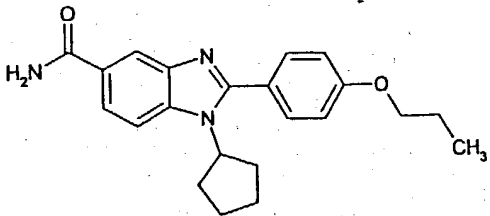
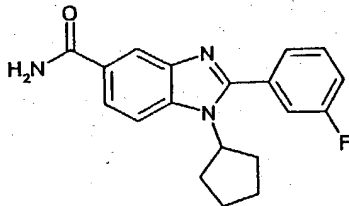
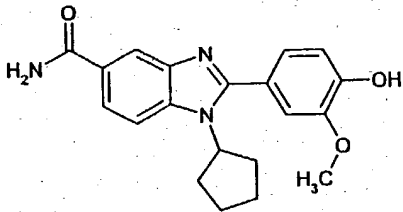
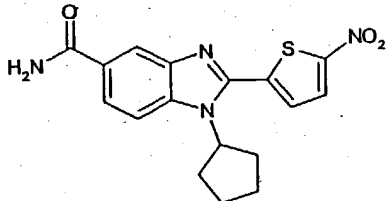
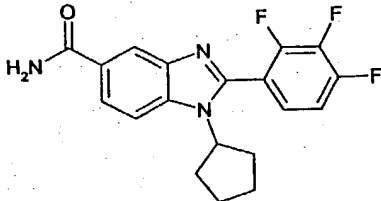
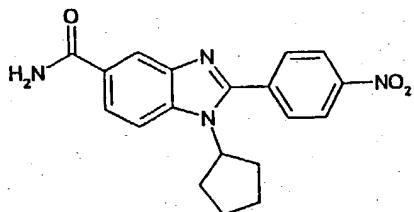
Ex. No.	Formula	MS
1059		364 (M+H)
1060		324 (M+H)
1061		352 (M+H)
1062		357 (M+H)
1063		360 (M+H)
1064		351 (M+H)

Table 82

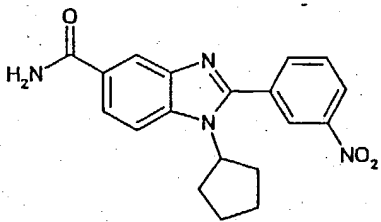
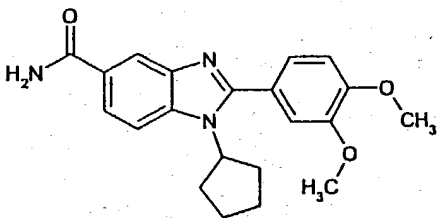
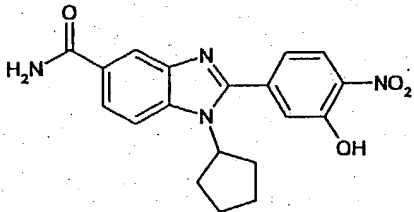
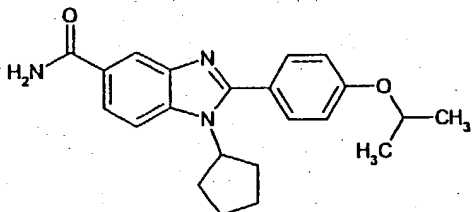
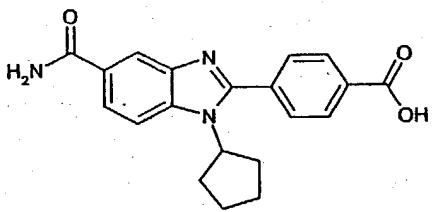
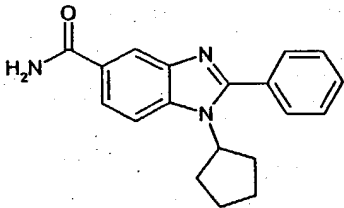
Ex. No.	Formula	MS
1065		351 (M+H)
1066		366 (M+H)
1067		367 (M+H)
1068		364 (M+H)
1069		350 (M+H)
1070		306 (M+H)

Table 83

Ex. No.	Formula	MS
1071		365 (M+H)
1072		455 (M+H)
1073		399 (M+H)
1074		358 (M+H)
1075		337 (M+H)
1076		386 (M+H)

Table 84

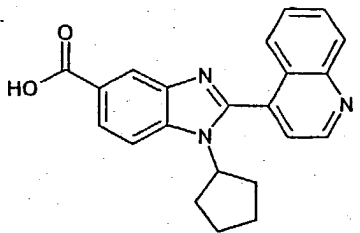
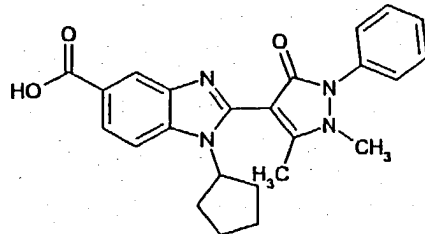
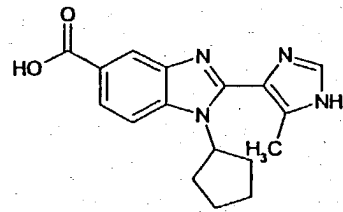
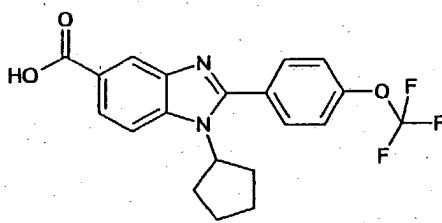
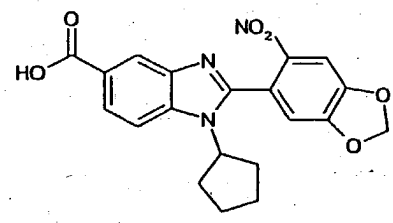
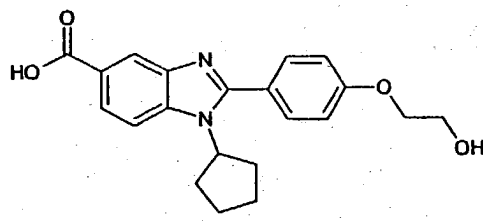
Ex. No.	Formula	MS
1077		358 (M+H)
1078		417 (M+H)
1079		311 (M+H)
1080		391 (M+H)
1081		396 (M+H)
1082		367 (M+H)



Table 85

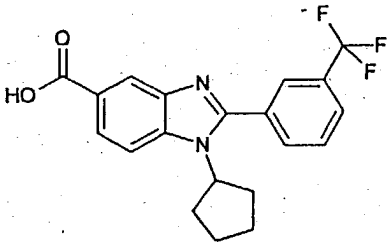
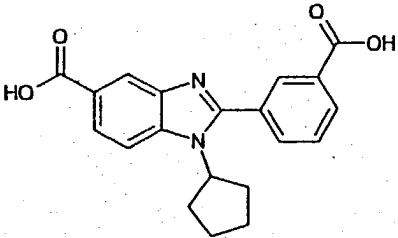
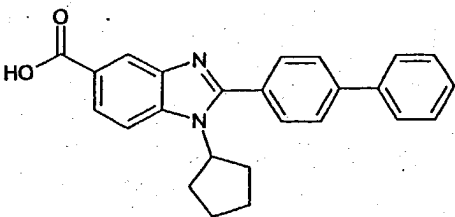
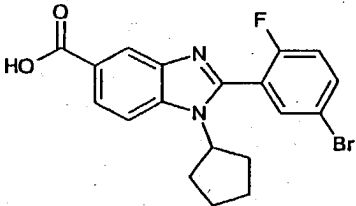
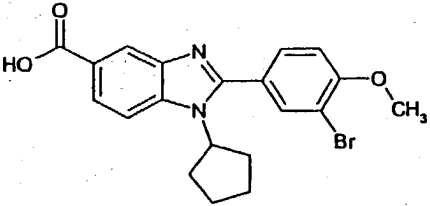
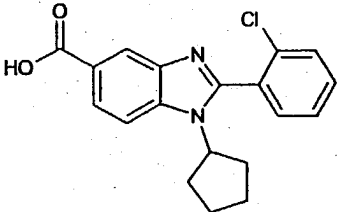
Ex. No.	Formula	MS
1083		375 (M+H)
1084		351 (M+H)
1085		383 (M+H)
1086		403 (M+H)
1087		415 (M+H)
1088		341 (M+H)

Table 86

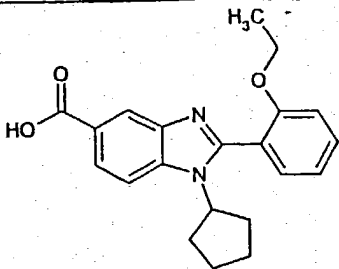
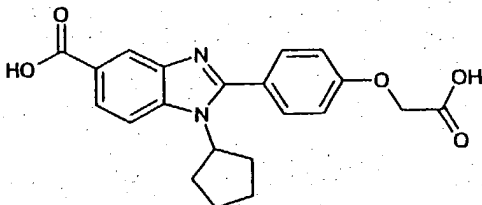
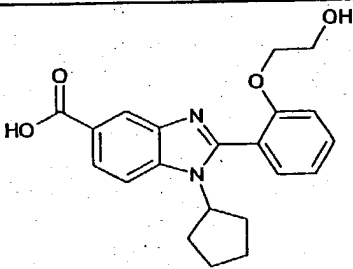
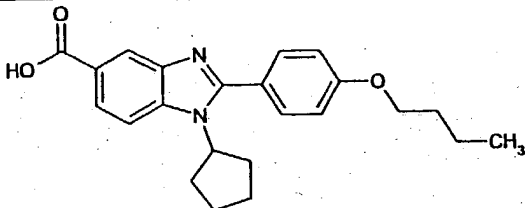
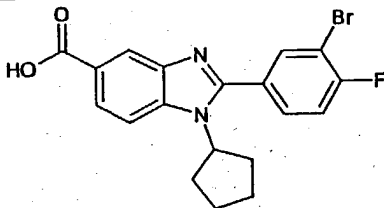
Ex. No.	Formula	MS
1089		351 (M+H)
1090		381 (M+H)
1091		367 (M+H)
1092		379 (M+H)
1093		403 (M+H)

Table 87

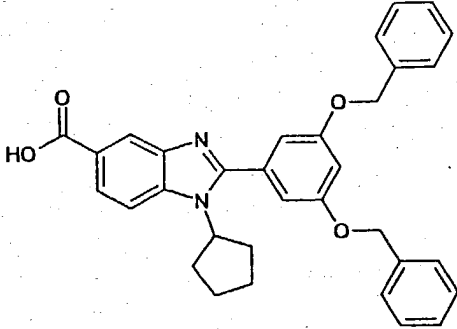
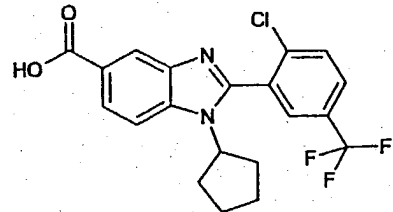
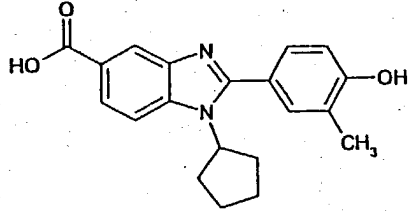
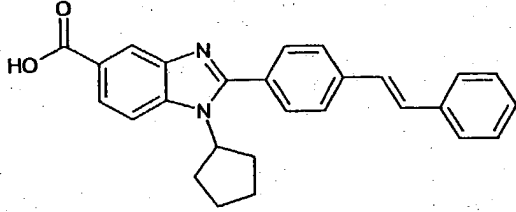
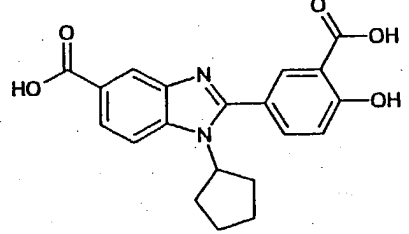
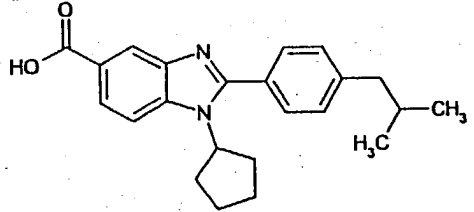
Ex. No.	Formula	MS
1094		519 (M+H)
1095		409 (M+H)
1096		337 (M+H)
1097		409 (M+H)
1098		367 (M+H)
1099		363 (M+H)

Table 88

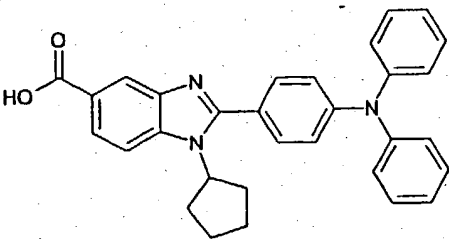
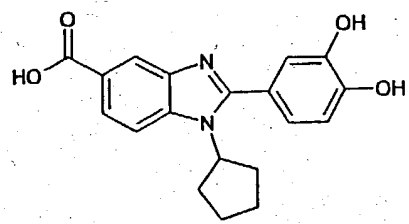
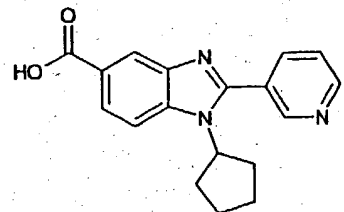
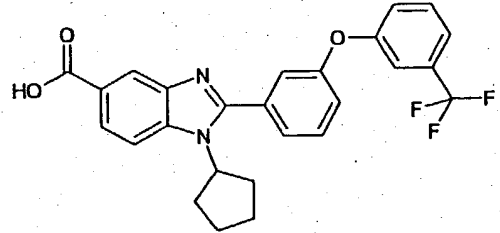
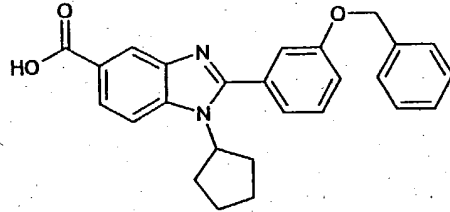
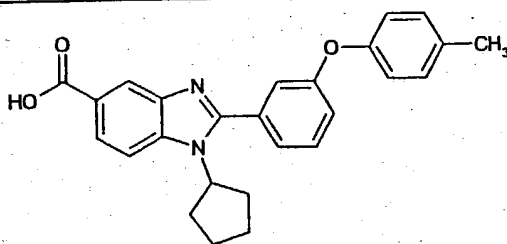
Ex. No.	Formula	MS
1100		474 (M+H)
1101		339 (M+H)
1102		308 (M+H)
1103		467 (M+H)
1104		413 (M+H)
1105		413 (M+H)

Table 89

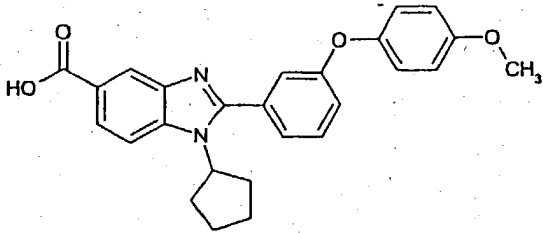
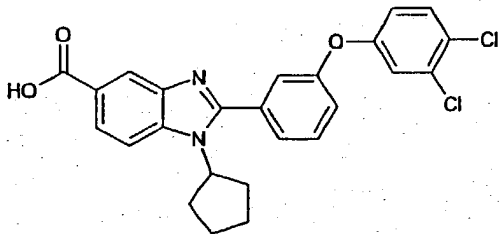
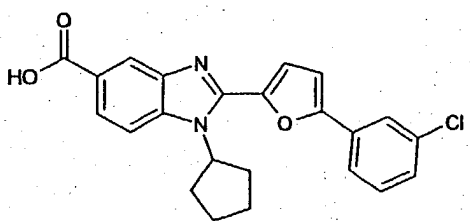
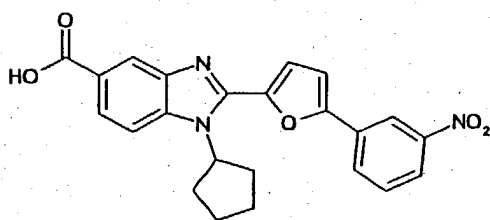
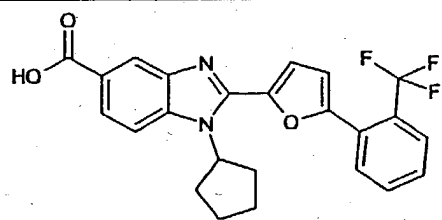
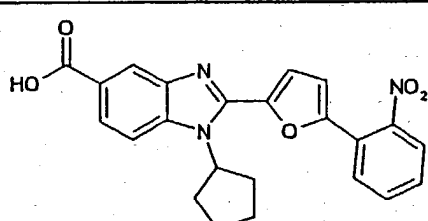
Ex. No.	Formula	MS
1106		429 (M+H)
1107		467 (M+H)
1108		
1109		
1110		441 (M+H)
1111		418 (M+H)

Table 90

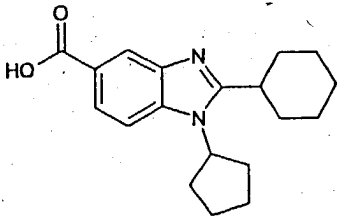
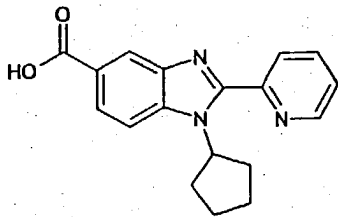
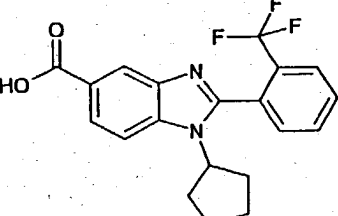
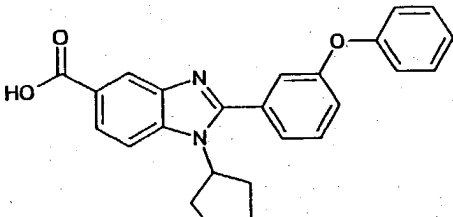
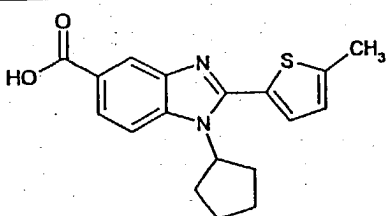
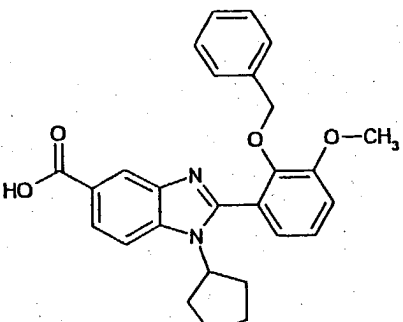
Ex. No.	Formula	MS
1112		313 (M+H)
1113		308 (M+H)
1114		375 (M+H)
1115		399 (M+H)
1116		327 (M+H)
1117		443 (M+H)

Table 91

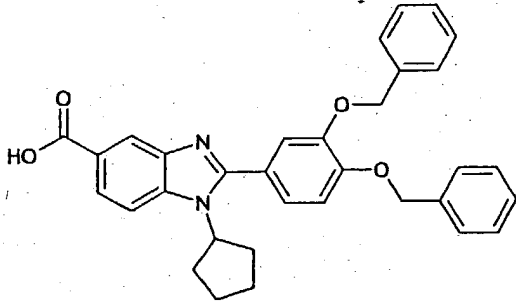
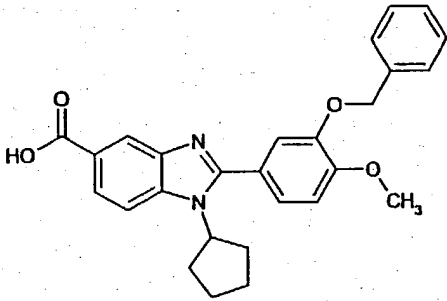
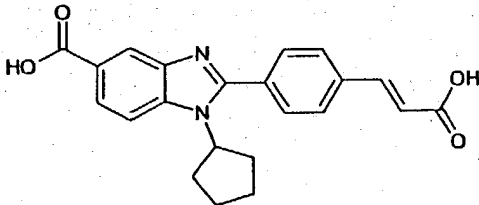
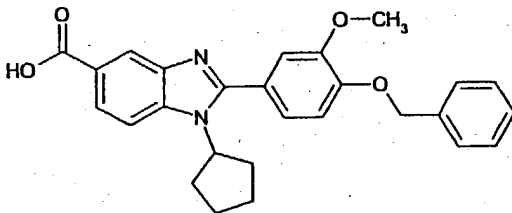
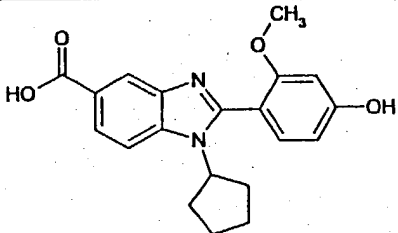
Ex. No.	Formula	MS
1118		519 (M+H)
1119		443 (M+H)
1120		377 (M+H)
1121		443 (M+H)
1122		353 (M+H)

Table 92

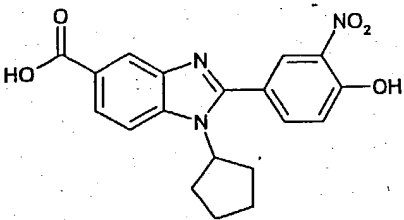
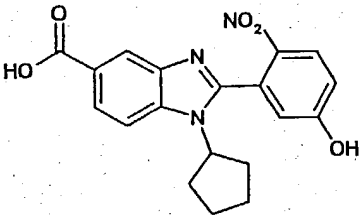
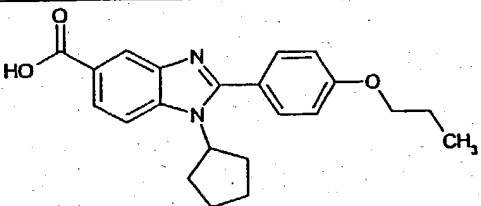
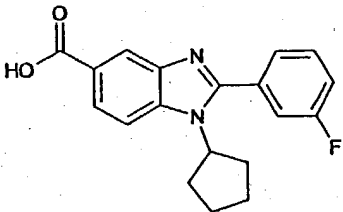
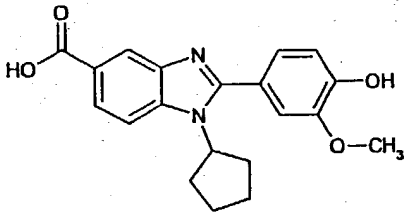
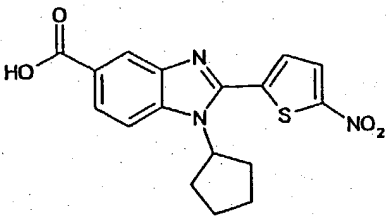
Ex. No.	Formula	MS
1123		368 (M+H)
1124		368 (M+H)
1125		365 (M+H)
1126		325 (M+H)
1127		353 (M+H)
1128		358 (M+H)



Table 93

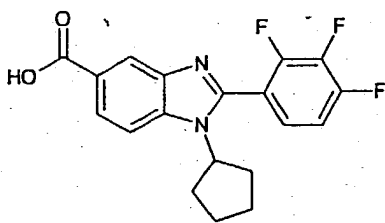
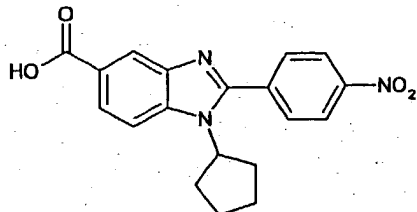
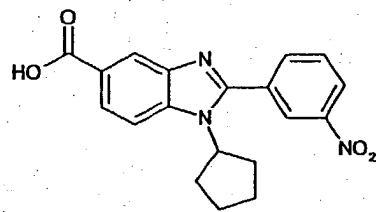
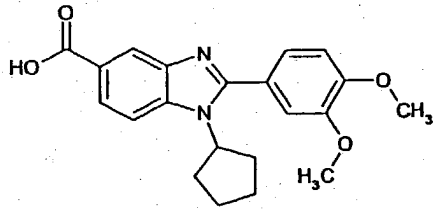
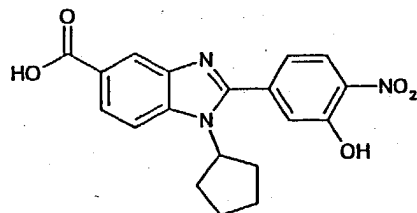
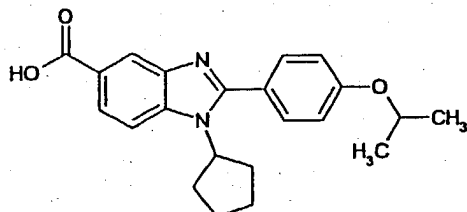
Ex. No.	Formula	MS
1129		361 (M+H)
1130		352 (M+H)
1131		352 (M+H)
1132		367 (M+H)
1133		368 (M+H)
1134		365 (M+H)

Table 94

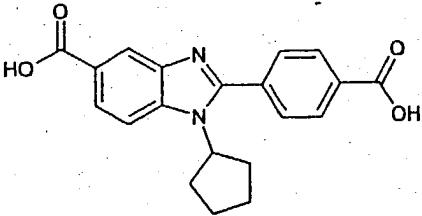
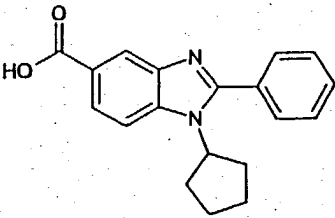
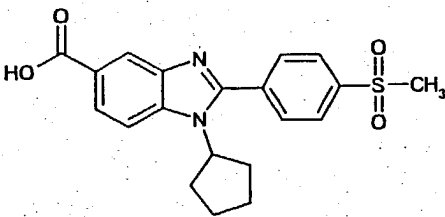
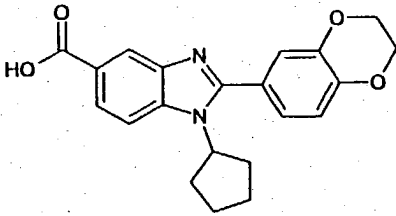
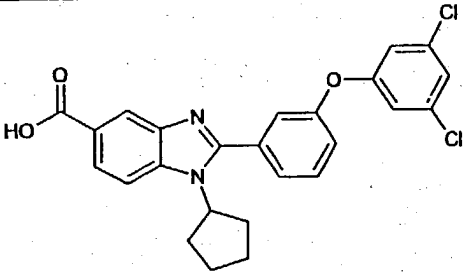
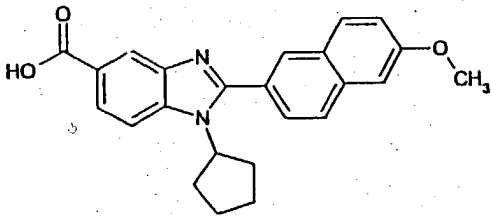
Ex. No.	Formula	MS
1135		351 (M+H)
1136		307 (M+H)
1137		385 (M+H)
1138		365 (M+H)
1139		467 (M+H)
1140		387 (M+H)

Table 95

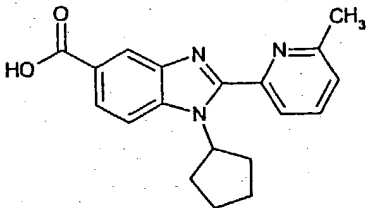
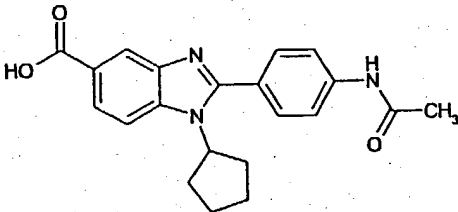
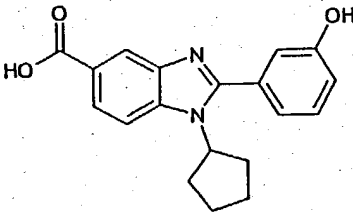
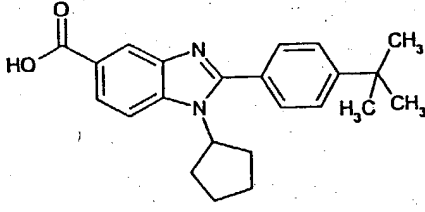
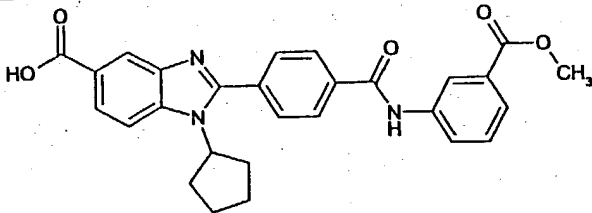
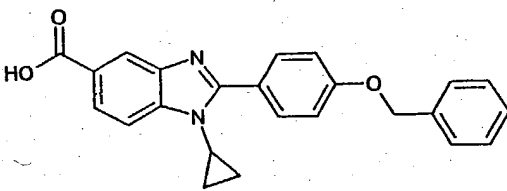
Ex. No.	Formula	MS
1141		322 (M+H)
1142		364 (M+H)
1143		323 (M+H)
1144		363 (M+H)
1145		484 (M+H)
1146		385 (M+H)

Table 96

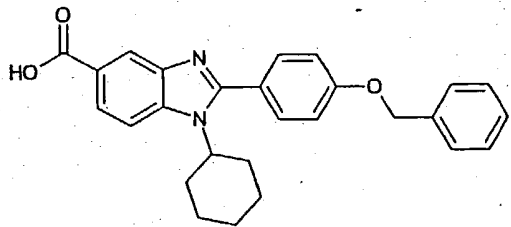
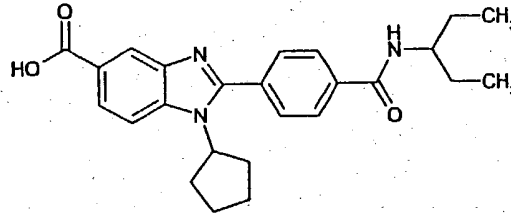
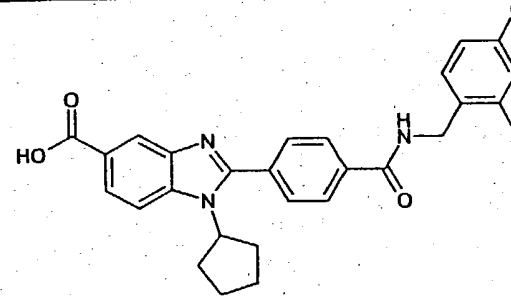
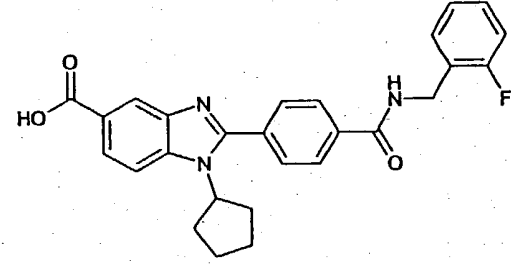
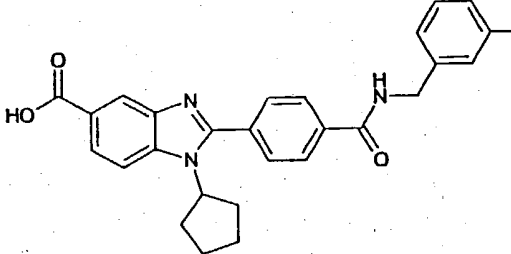
Ex. No.	Formula	MS
1147		427 (M+H)
1148		420 (M+H)
1149		508 (M+H)
1150		458 (M+H)
1151		458 (M+H)

Table 97

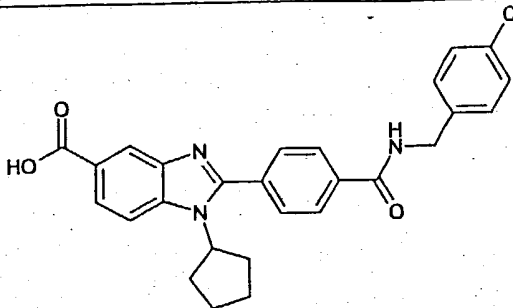
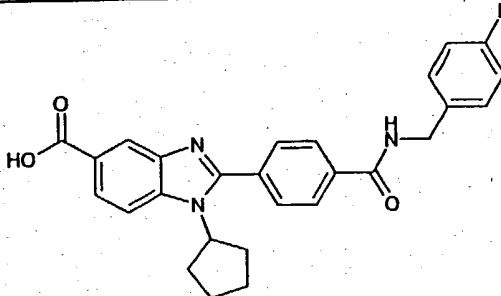
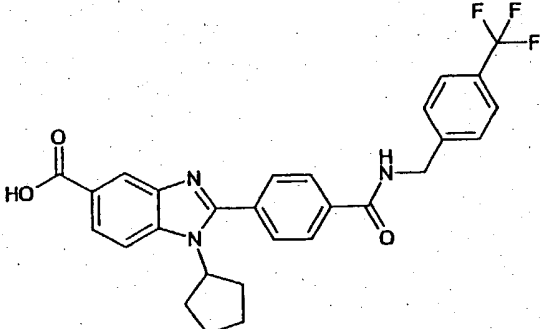
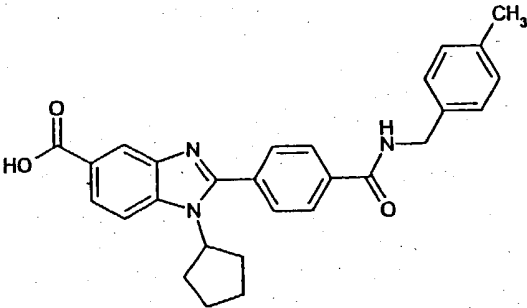
Ex. No.	Formula	MS
1152		474 (M+H)
1153		458 (M+H)
1154		508 (M+H)
1155		454 (M+H)

Table 98

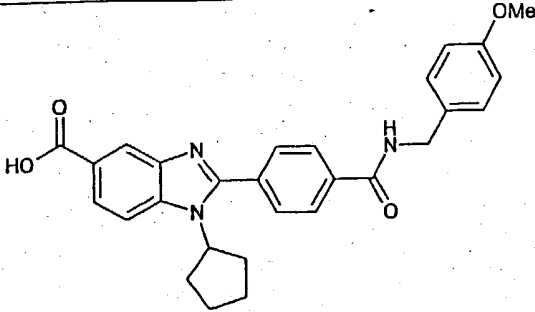
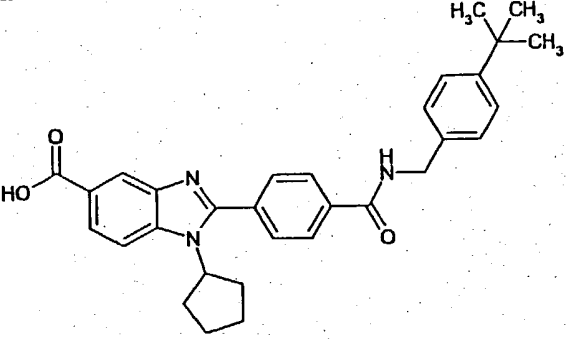
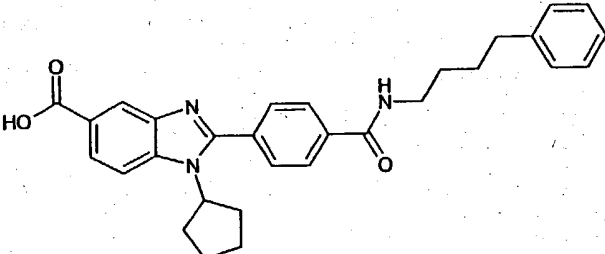
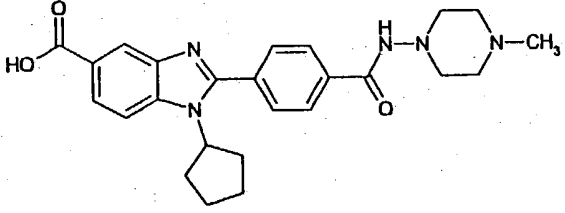
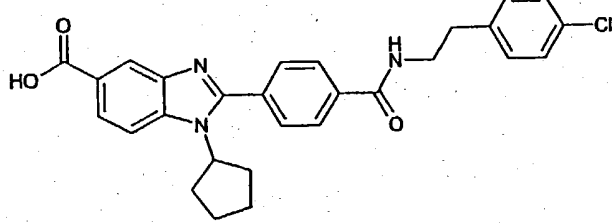
Ex. No.	Formula	MS
1156		470 (M+H)
1157		496 (M+H)
1158		482 (M+H)
1159		448 (M+H)
1160		488 (M+H)

Table 99

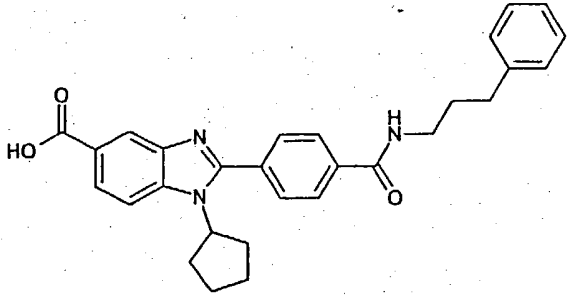
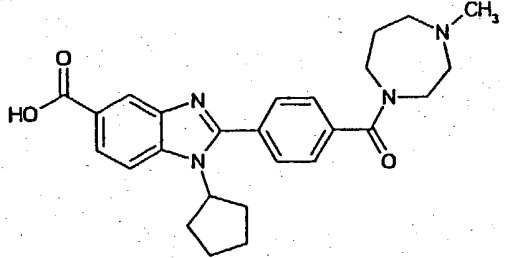
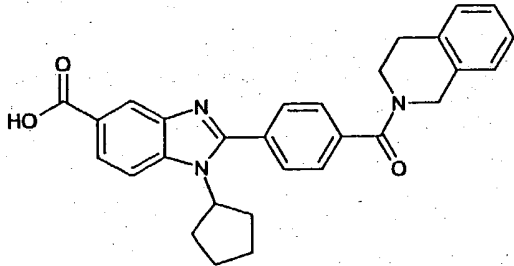
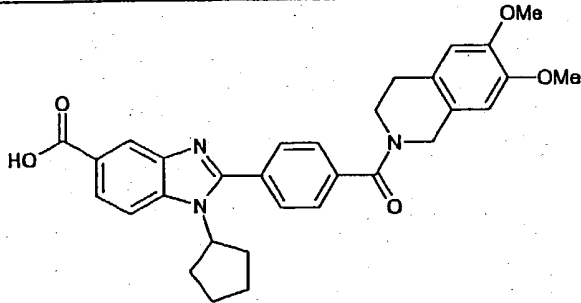
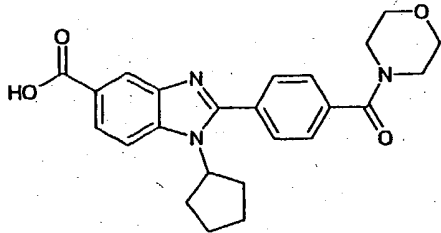
Ex. No.	Formula	MS
1161		468 (M+H)
1162		447 (M+H)
1163		466 (M+H)
1164		526 (M+H)
1165		420 (M+H)

Table 100

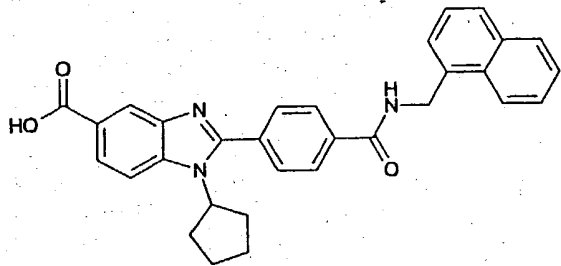
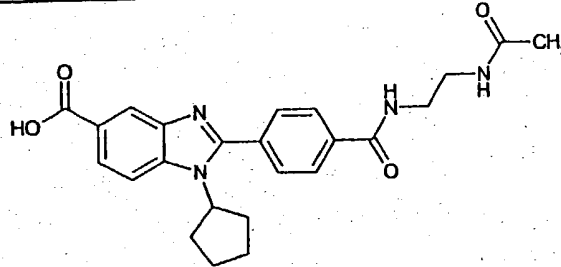
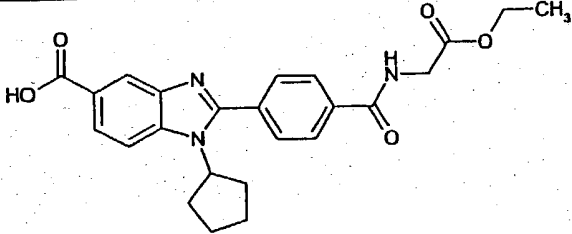
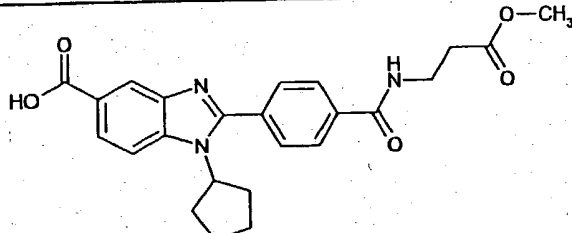
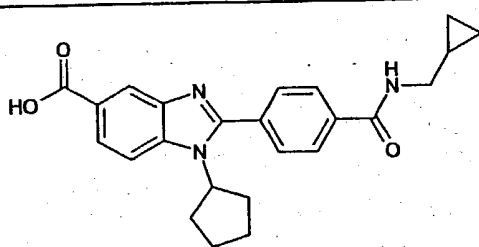
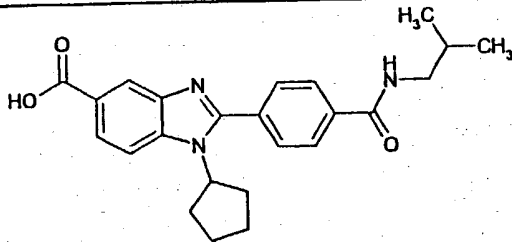
Ex. No.	Formula	MS
1166		490 (M+H)
1167		435 (M+H)
1168		436 (M+H)
1169		436 (M+H)
1170		404 (M+H)
1171		406 (M+H)



Table 101

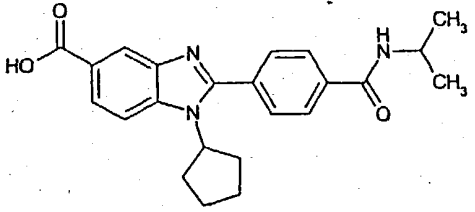
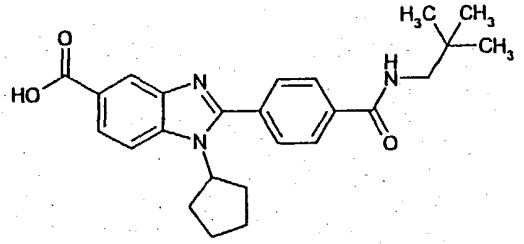
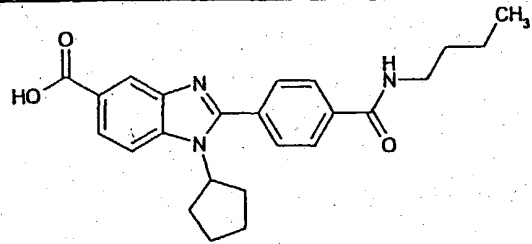
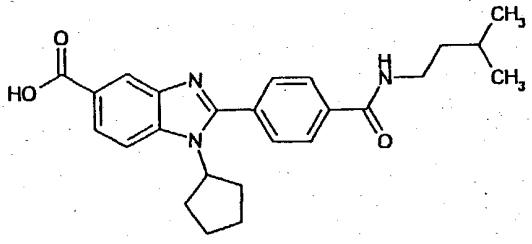
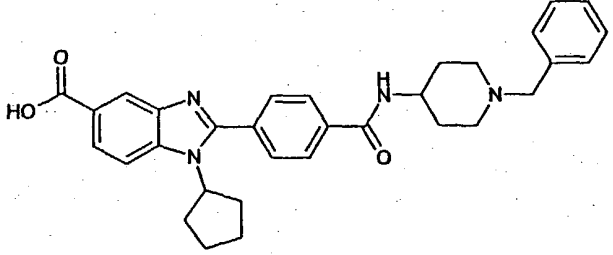
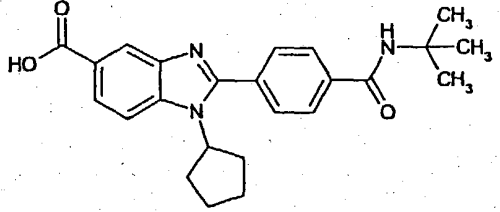
Ex. No.	Formula	MS
1172		392 (M+H)
1173		420 (M+H)
1174		406 (M+H)
1175		420 (M+H)
1176		523 (M+H)
1177		406 (M+H)

Table 102

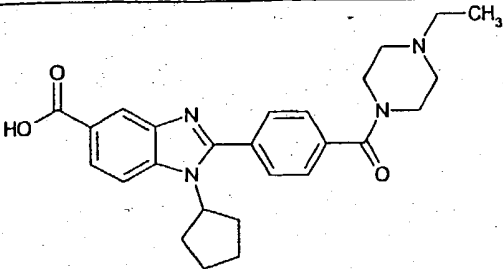
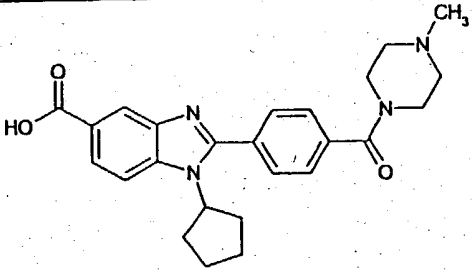
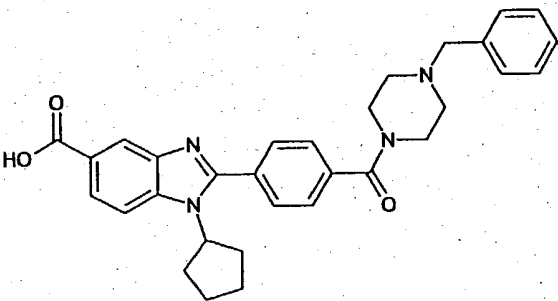
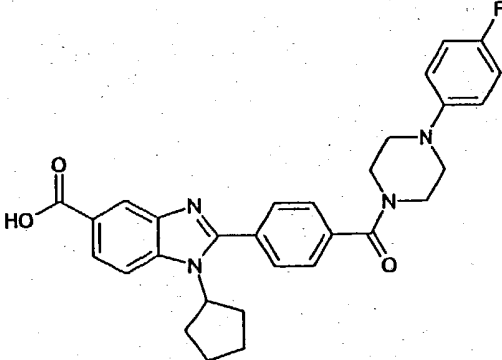
Ex. No.	Formula	MS
1178		447 (M+H)
1179		433 (M+H)
1180		509 (M+H)
1181		513 (M+H)

Table 103

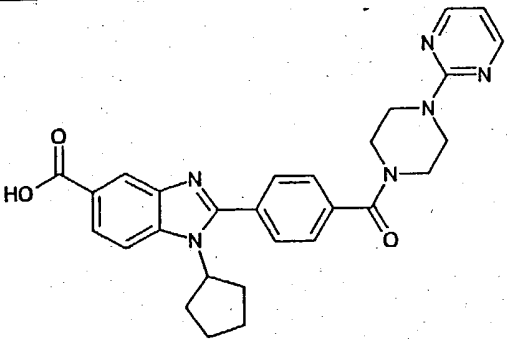
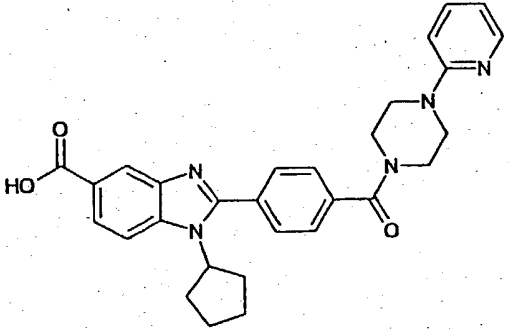
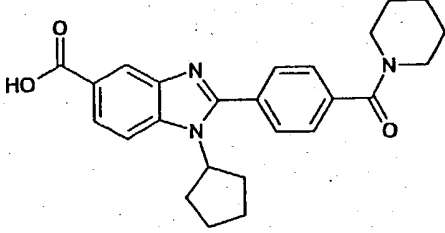
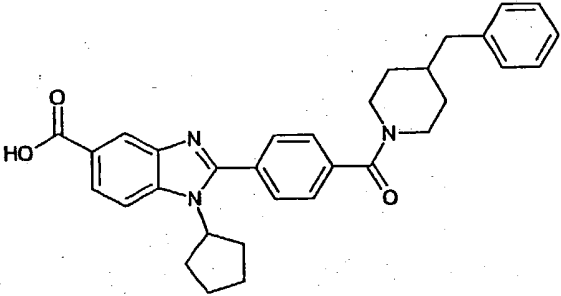
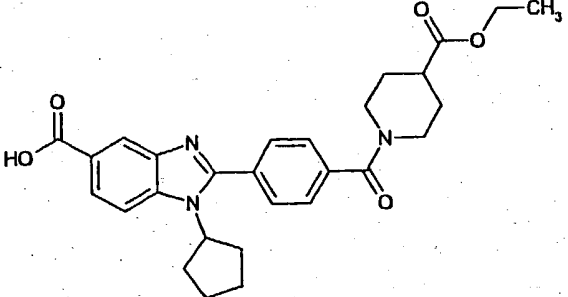
Ex. No.	Formula	MS
1182		497 (M+H)
1183		496 (M+H)
1184		418 (M+H)
1185		508 (M+H)
1186		490 (M+H)

Table 104

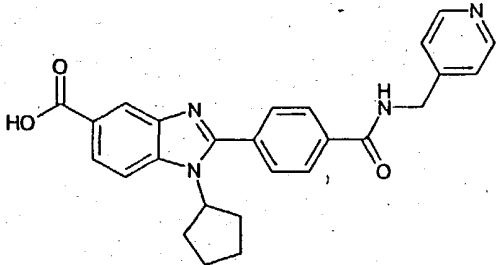
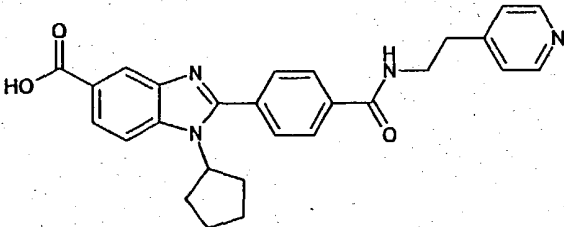
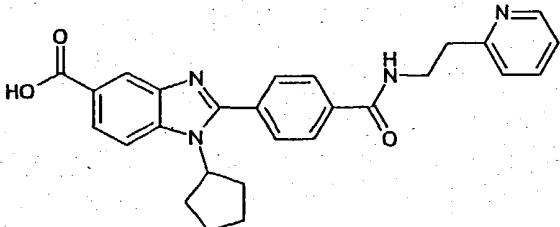
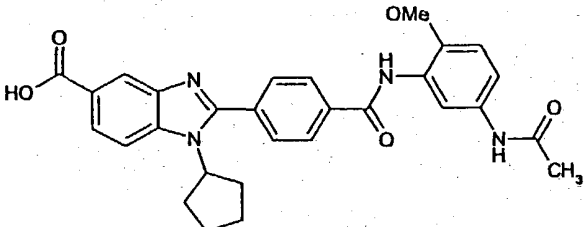
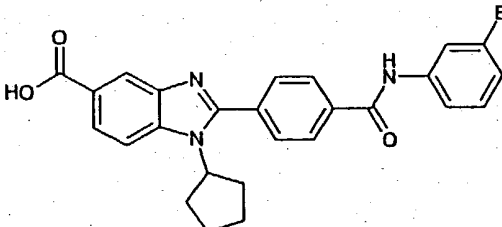
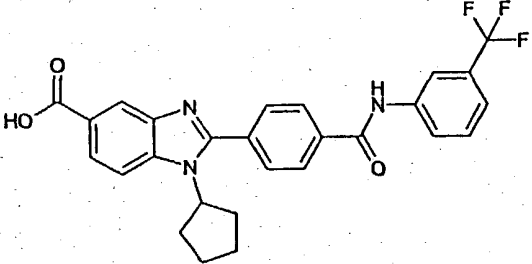
Ex. No.	Formula	MS
1187		441 (M+H)
1188		455 (M+H)
1189		455 (M+H)
1190		513 (M+H)
1191		504 (M+H)
1192		494 (M+H)

Table 105

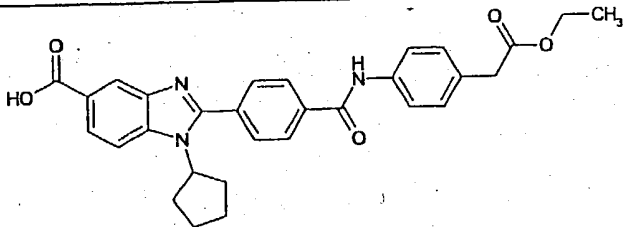
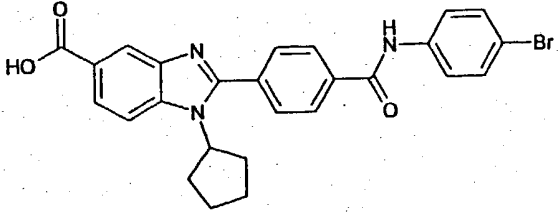
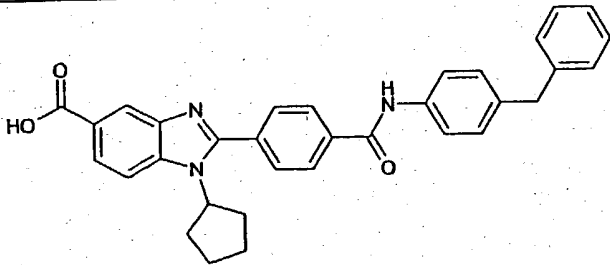
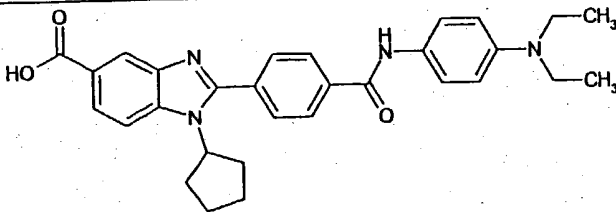
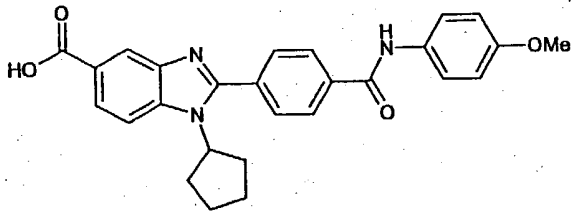
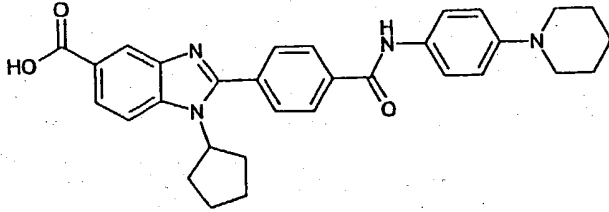
Ex. No.	Formula	MS
1193		512 (M+H)
1194		504 (M+H)
1195		516 (M+H)
1196		497 (M+H)
1197		456 (M+H)
1198		509 (M+H)

Table 106

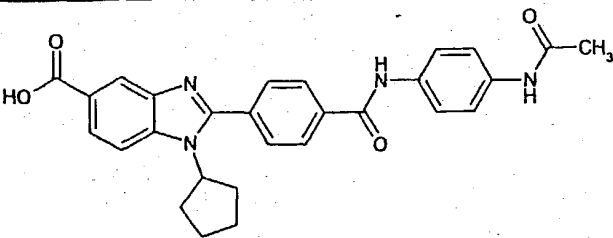
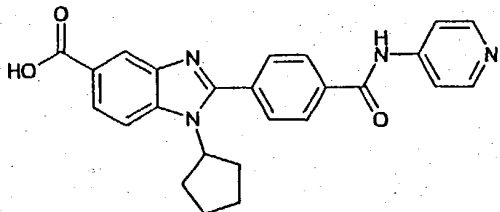
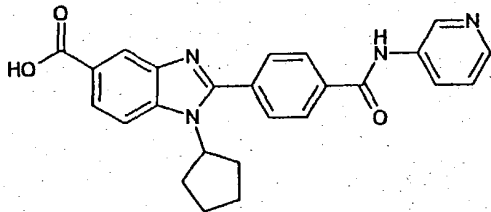
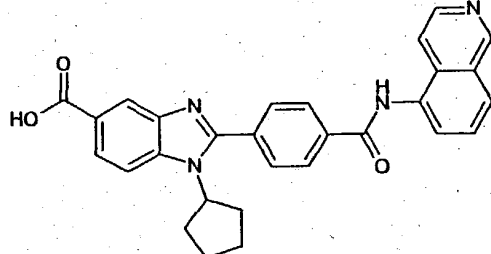
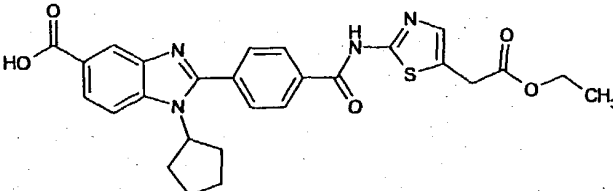
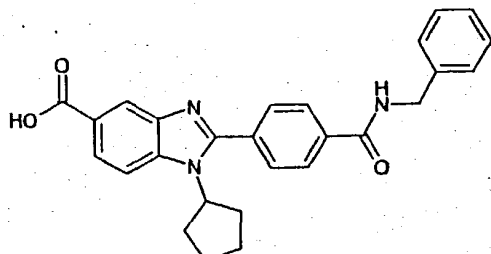
Ex. No.	Formula	MS
1199		483 (M+H)
1200		427 (M+H)
1201		427 (M+H)
1202		477 (M+H)
1203		519 (M+H)
1204		440 (M+H)

Table 107

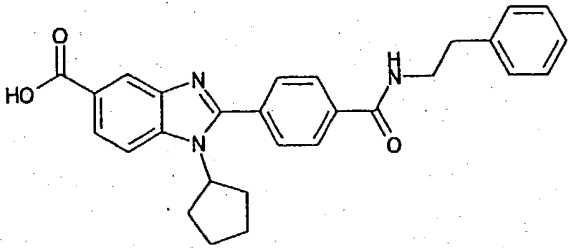
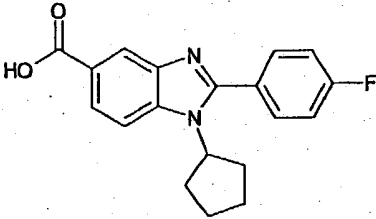
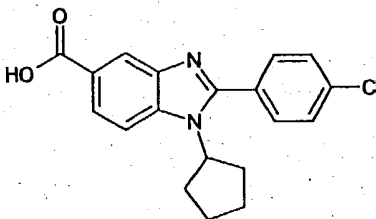
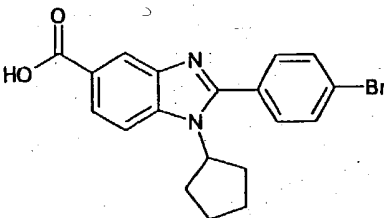
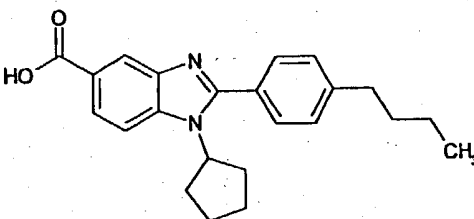
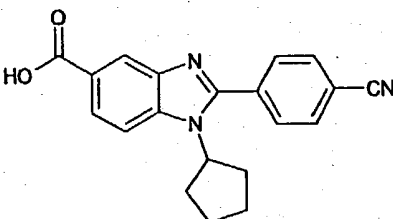
Ex. No.	Formula	MS
1205		454 (M+H)
1206		325 (M+H)
1207		341 (M+H)
1208		385 (M+H)
1209		363 (M+H)
1210		332 (M+H)

Table 108

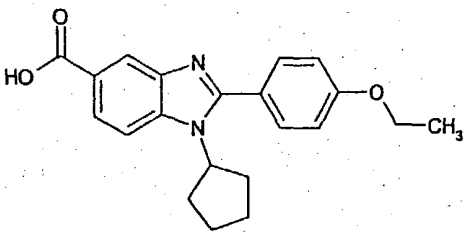
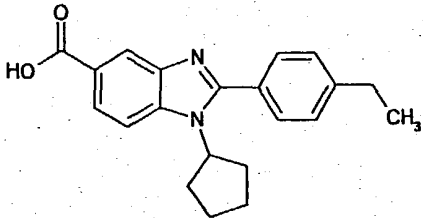
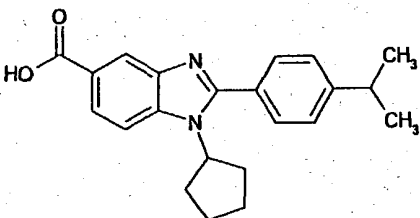
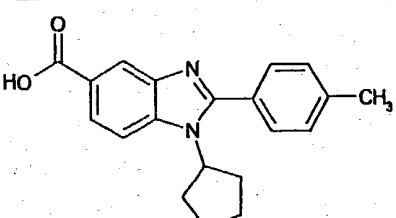
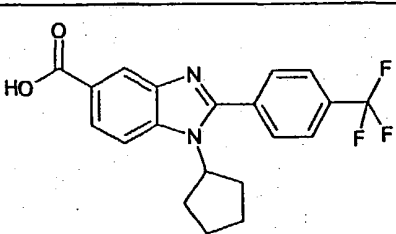
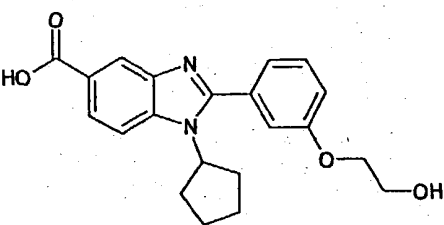
Ex. No.	Formula	MS
1211		351 (M+H)
1212		335 (M+H)
1213		349 (M+H)
1214		321 (M+H)
1215		375 (M+H)
1216		367 (M+H)



Table 109

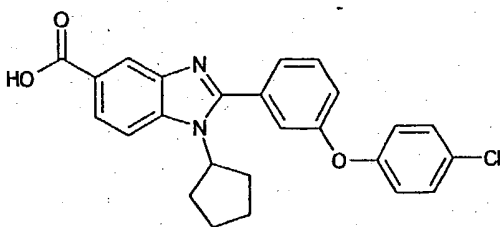
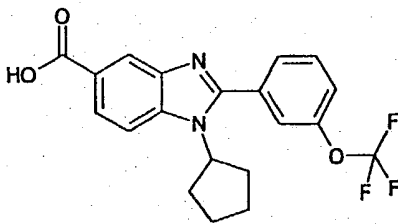
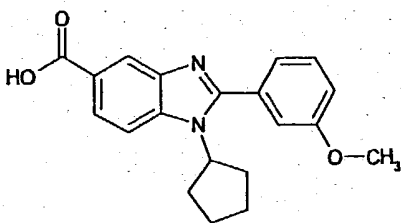
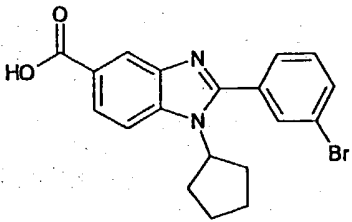
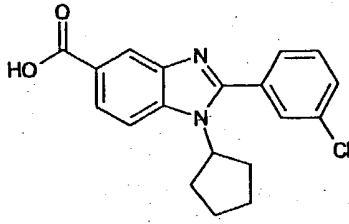
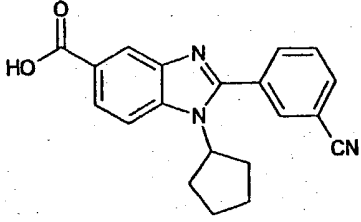
Ex. No.	Formula	MS
1217		433 (M+H)
1218		391 (M+H)
1219		337 (M+H)
1220		385 (M+H)
1221		341 (M+H)
1222		332 (M+H)

Table 110

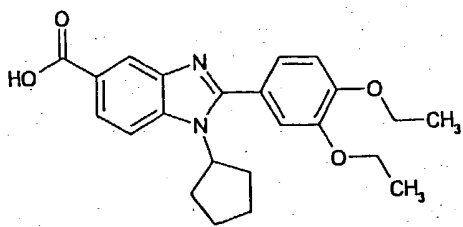
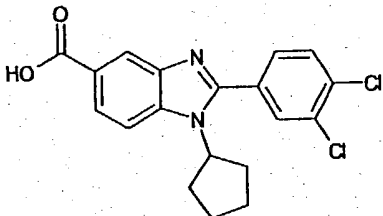
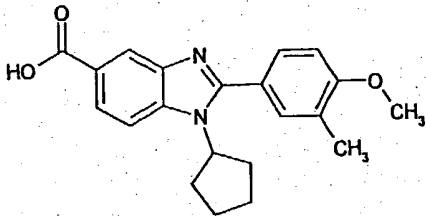
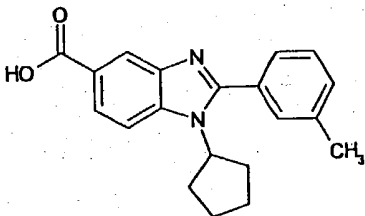
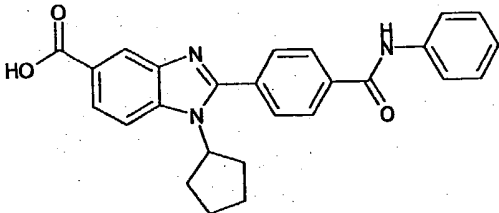
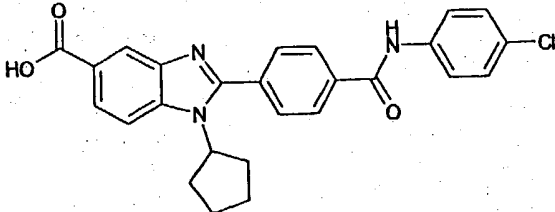
Ex. No.	Formula	MS
1223		395 (M+H)
1224		375 (M+H)
1225		351 (M+H)
1226		321 (M+H)
1227		426 (M+H)
1228		460 (M+H)

Table 111

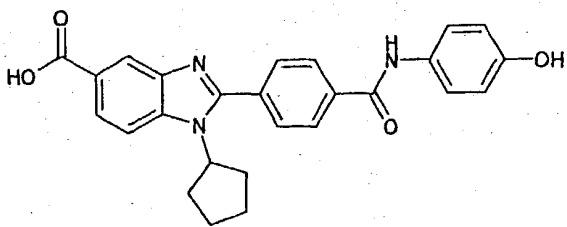
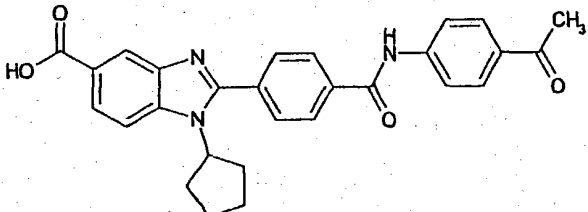
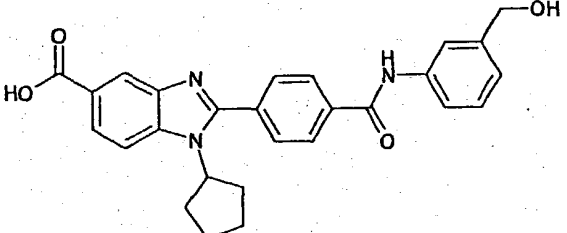
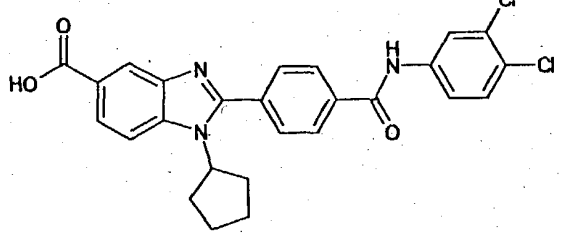
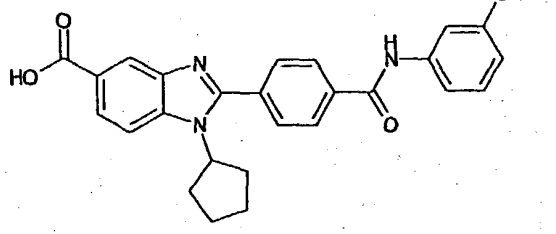
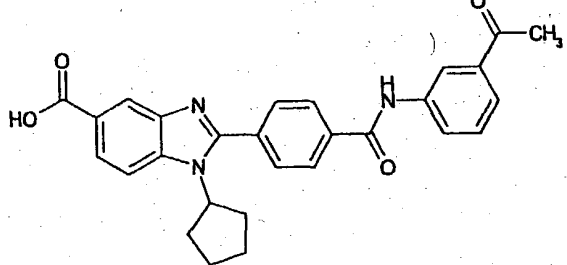
Ex. No.	Formula	MS
1229		442 (M+H)
1230		468 (M+H)
1231		456 (M+H)
1232		494 (M+H)
1233		451 (M+H)
1234		468 (M+H)

Table 112

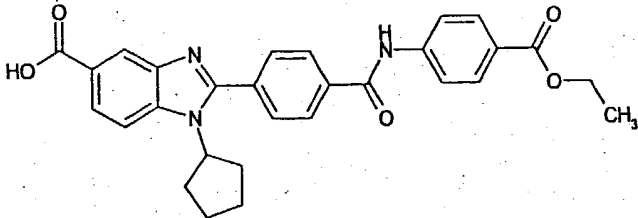
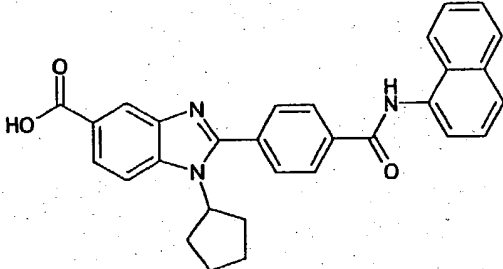
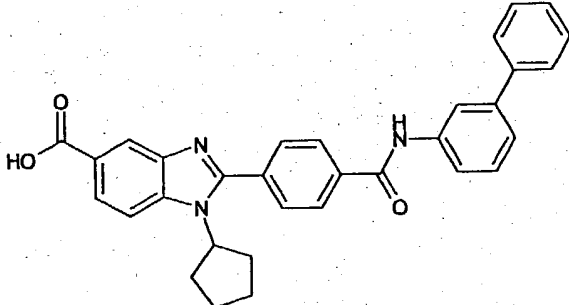
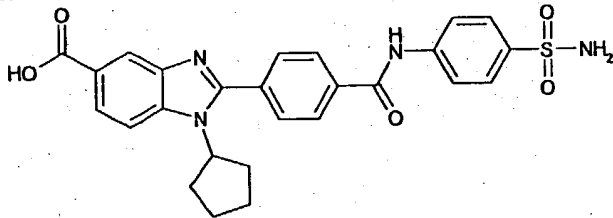
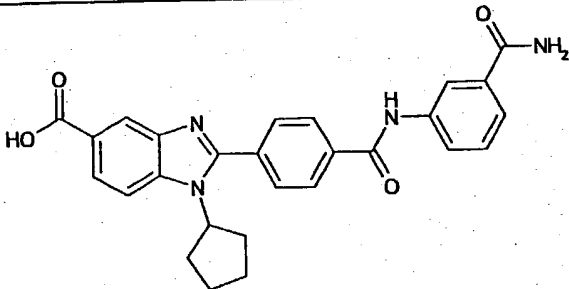
Ex. No.	Formula	MS
1235		498 (M+H)
1236		476 (M+H)
1237		502 (M+H)
1238		505 (M+H)
1239		469 (M+H)

Table 113

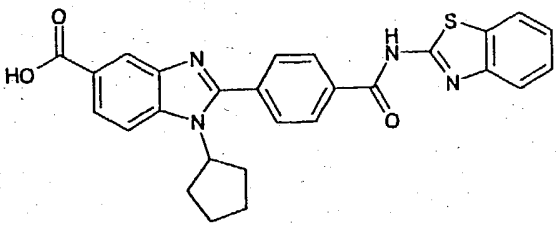
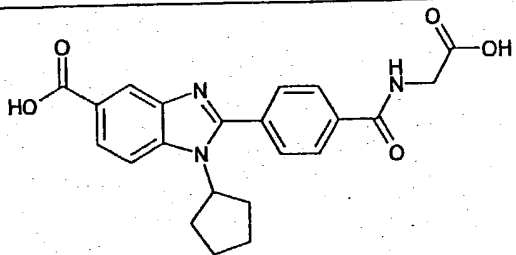
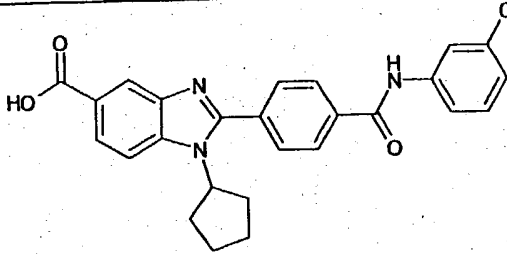
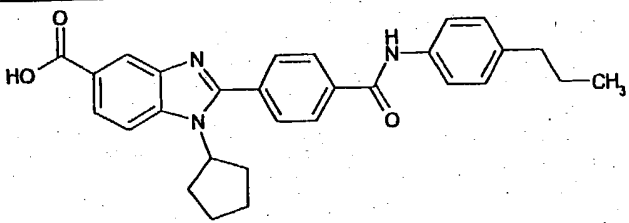
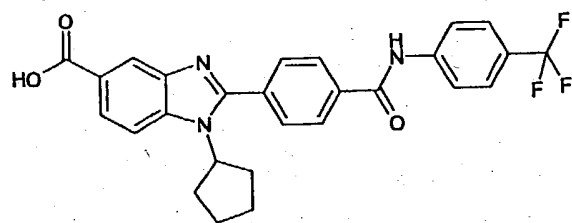
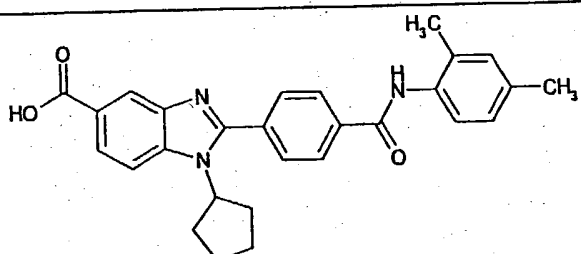
Ex. No.	Formula	MS
1240		483 (M+H)
1241		408 (M+H)
1242		460 (M+H)
1243		468 (M+H)
1244		494 (M+H)
1245		454 (M+H)

Table 114

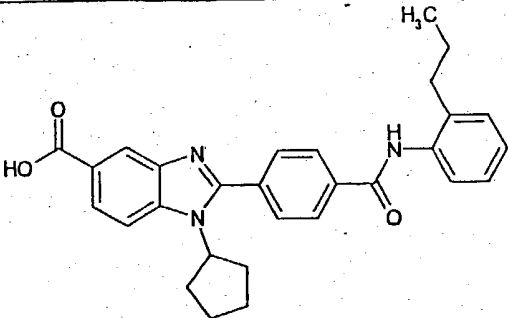
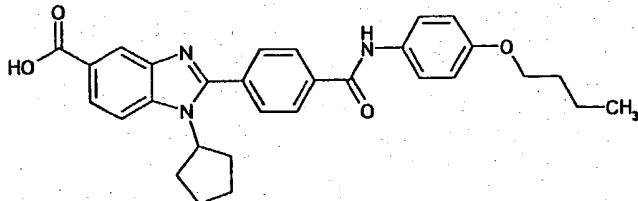
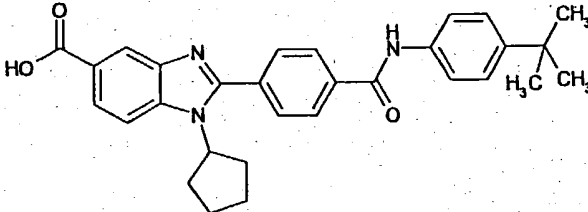
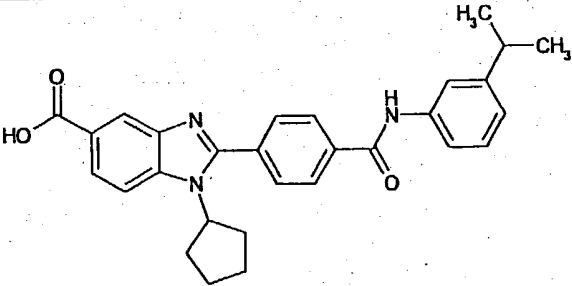
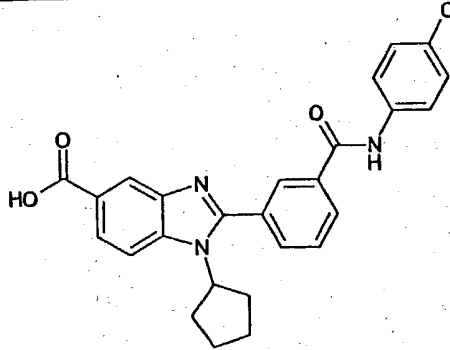
Ex. No.	Formula	MS
1246		468 (M+H)
1247		498 (M+H)
1248		482 (M+H)
1249		468 (M+H)
1250		460 (M+H)

Table 115

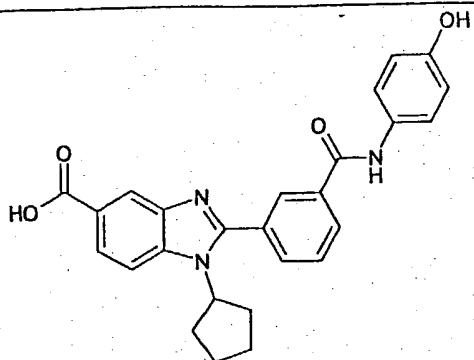
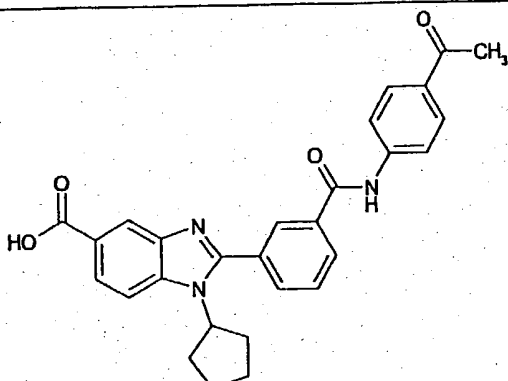
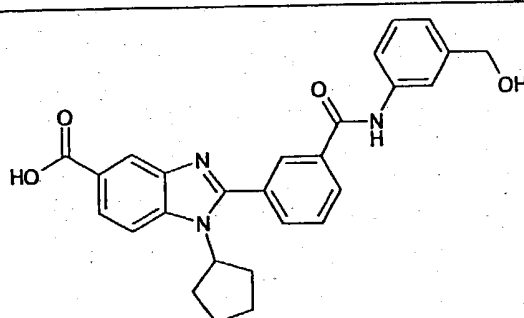
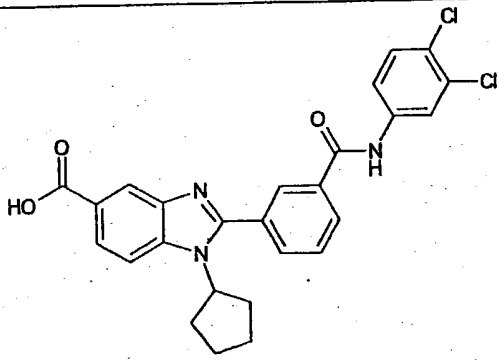
Ex. No.	Formula	MS
1251		442 (M+H)
1252		468 (M+H)
1253		456 (M+H)
1254		494 (M+H)

Table 116

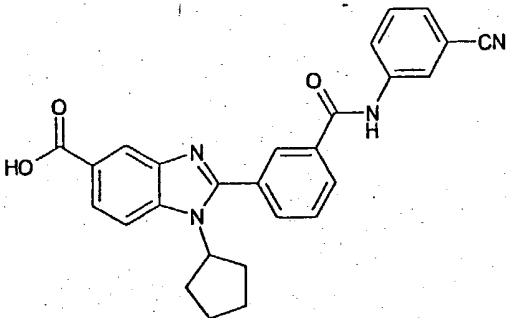
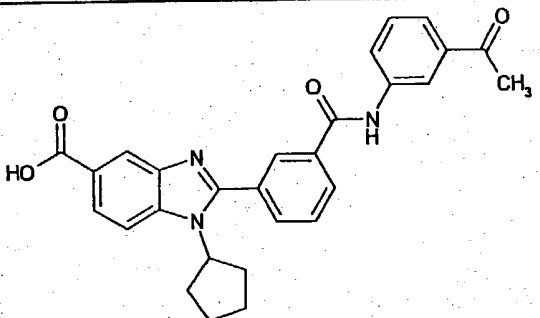
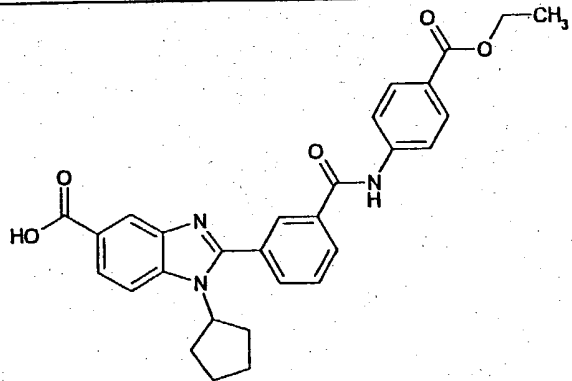
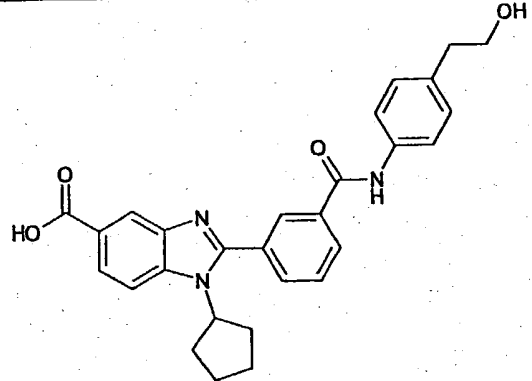
Ex. No.	Formula	MS
1255		451 (M+H)
1256		468 (M+H)
1257		498 (M+H)
1258		470 (M+H)



Table 117

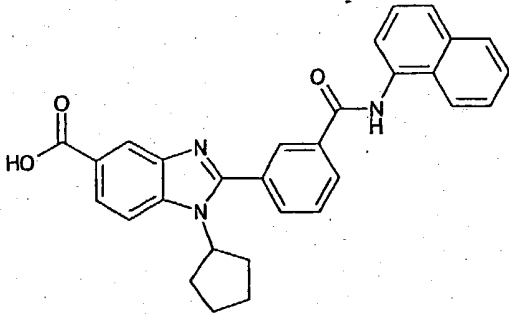
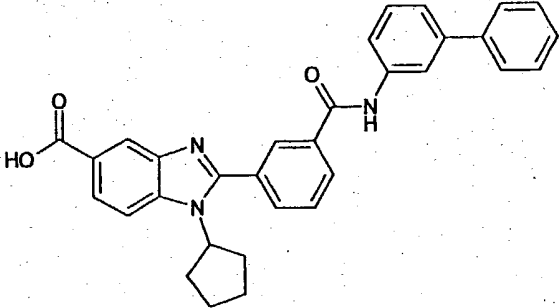
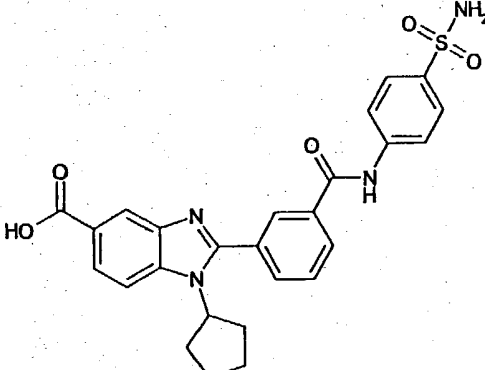
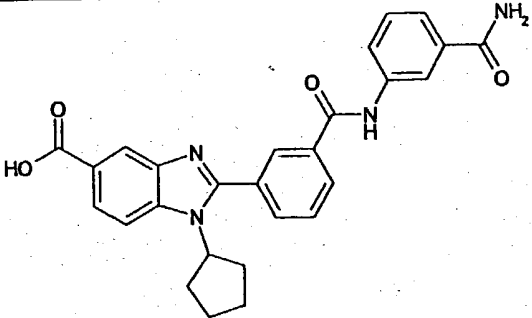
Ex. No.	Formula	MS
1259		476 (M+H)
1260		502 (M+H)
1261		505 (M+H)
1262		469 (M+H)

Table 118

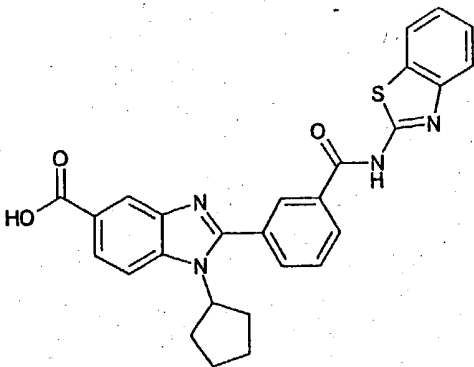
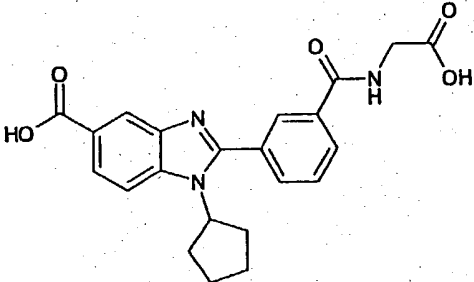
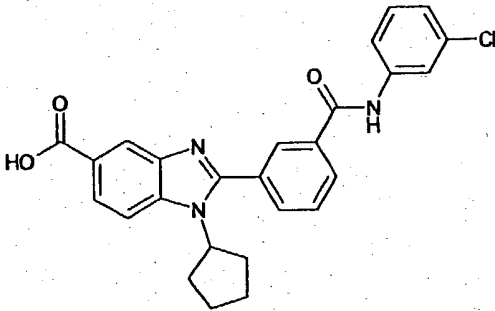
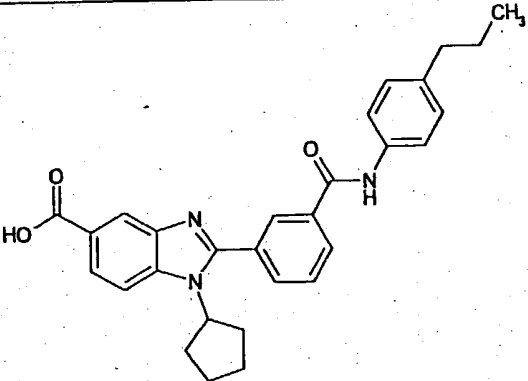
Ex. No.	Formula	MS
1263		483 (M+H)
1264		408 (M+H)
1265		460 (M+H)
1266		468 (M+H)

Table 119

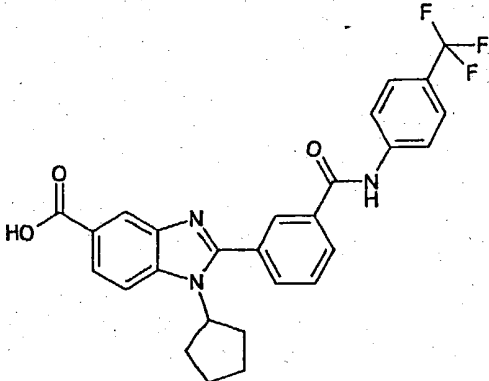
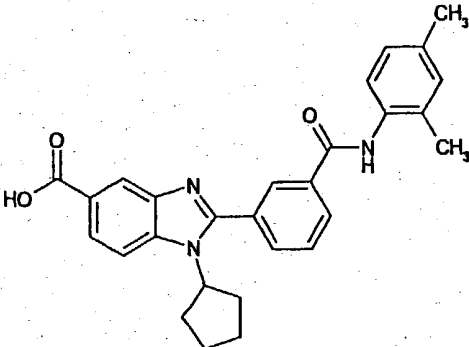
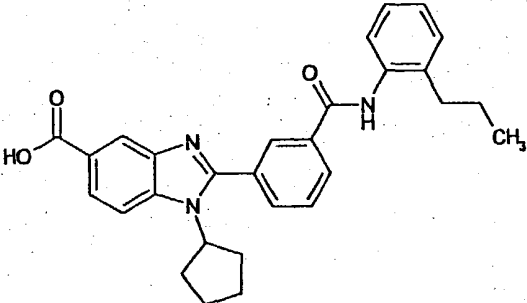
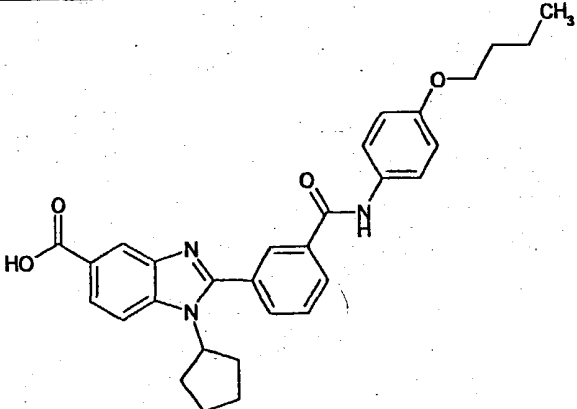
Ex. No.	Formula	MS
1267		494 (M+H)
1268		454 (M+H)
1269		468 (M+H)
1270		498 (M+H)

Table 120

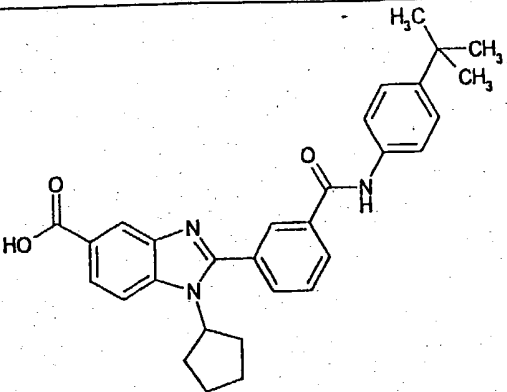
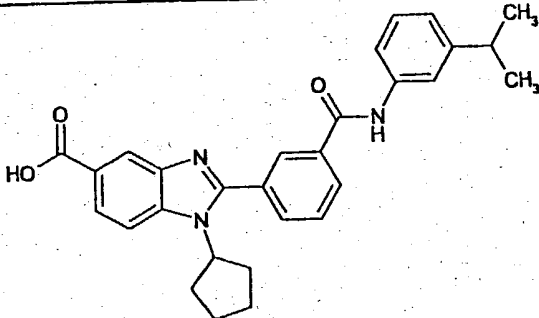
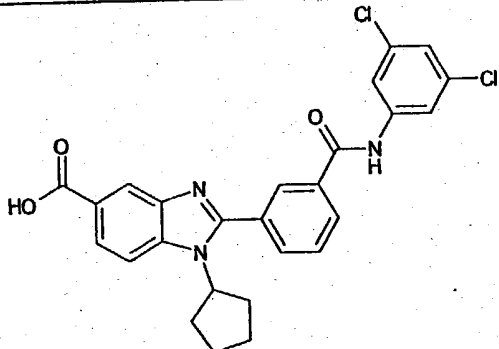
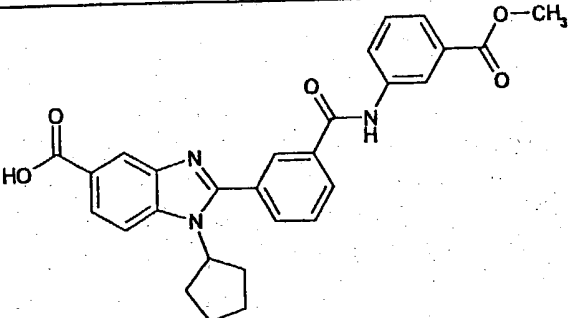
Ex. No.	Formula	MS
1271		482 (M+H)
1272		468 (M+H)
1273		494 (M+H)
1274		484 (M+H)

Table 121

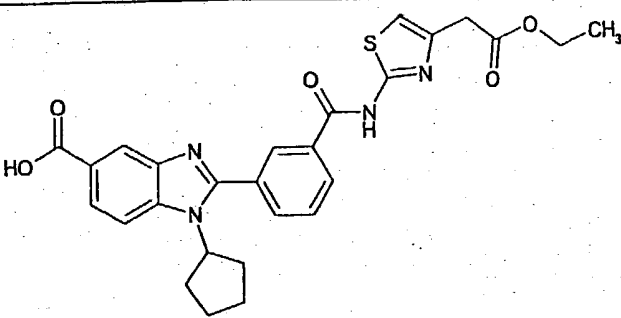
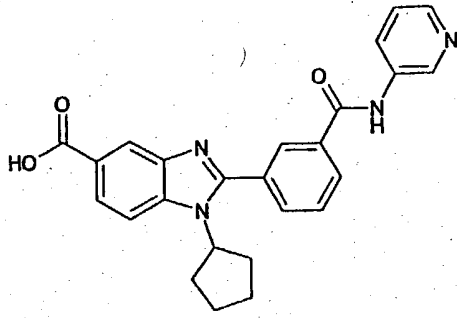
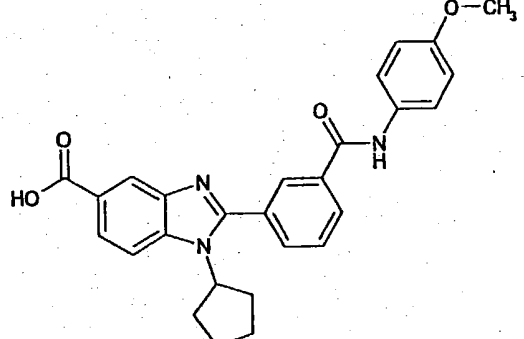
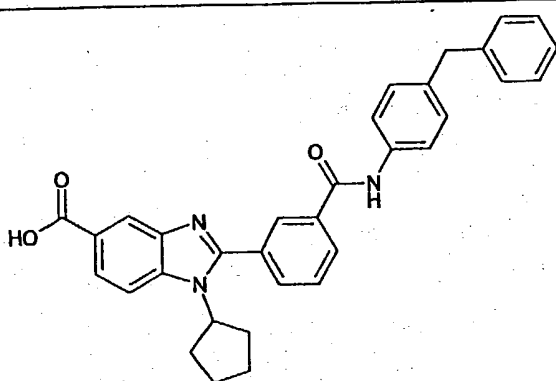
Ex. No.	Formula	MS
1275		519 (M+H)
1276		427 (M+H)
1277		456 (M+H)
1278		516 (M+H)

Table 122

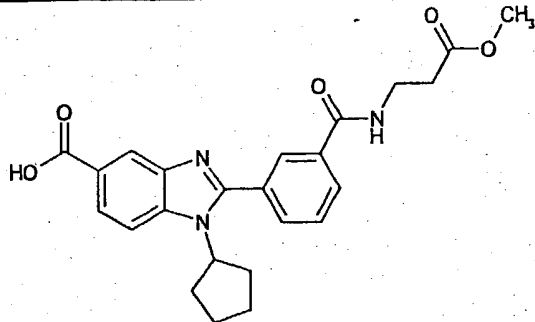
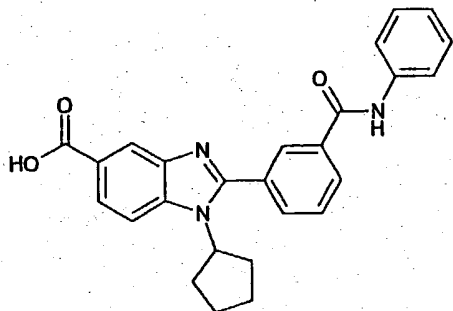
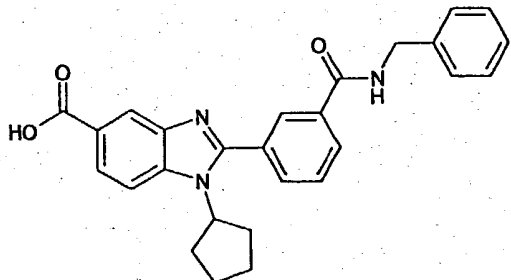
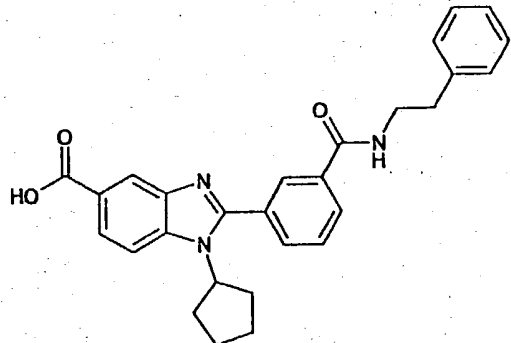
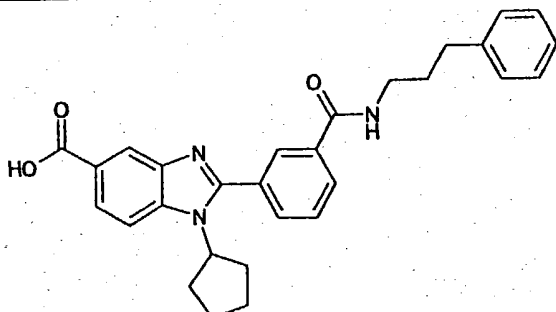
Ex. No.	Formula	MS
1279		436 (M+H)
1280		426 (M+H)
1281		440 (M+H)
1282		454 (M+H)
1283		468 (M+H)

Table 123

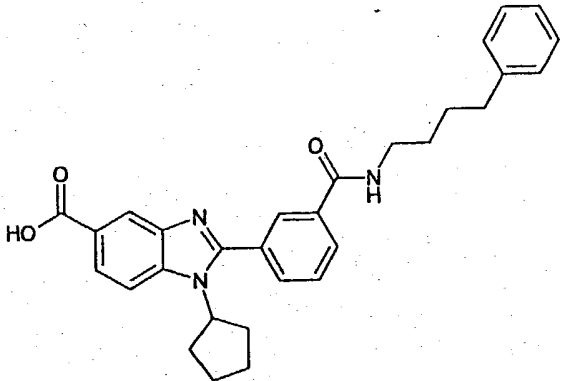
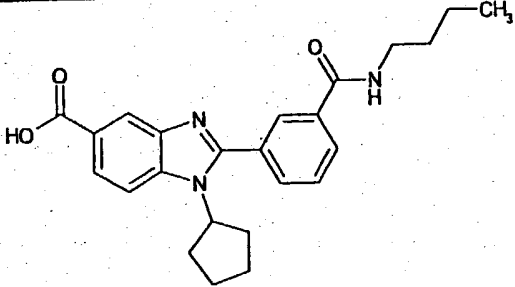
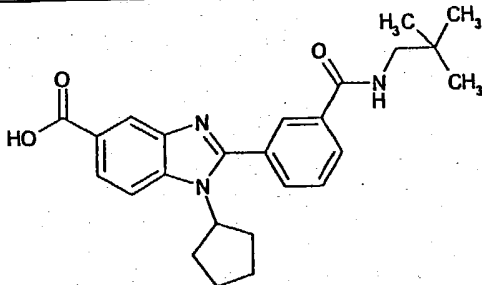
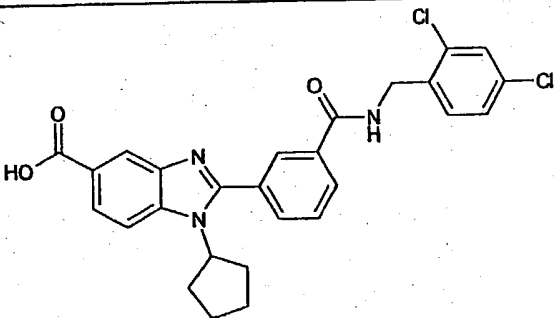
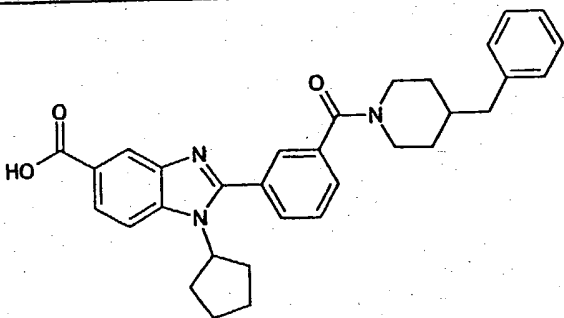
Ex. No.	Formula	MS
1284		482 (M+H)
1285		406 (M+H)
1286		420 (M+H)
1287		508 (M+H)
1288		508 (M+H)

Table 124

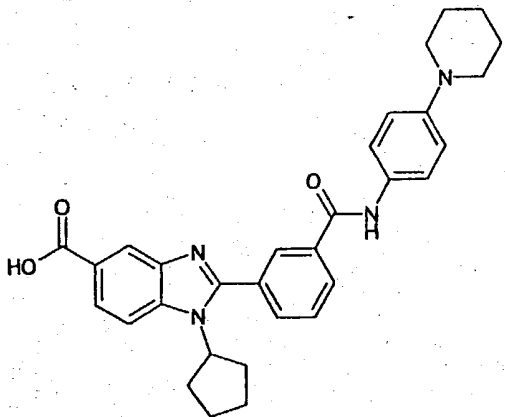
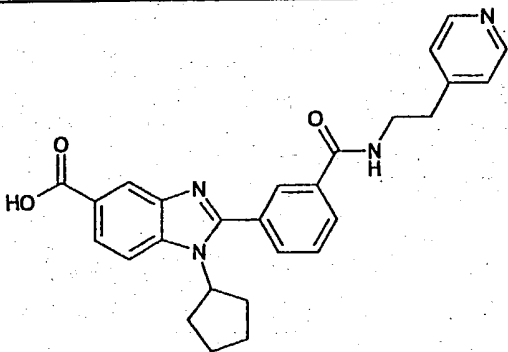
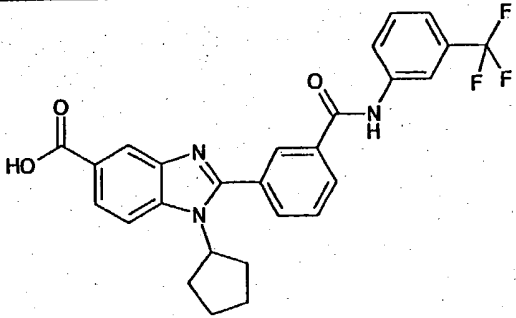
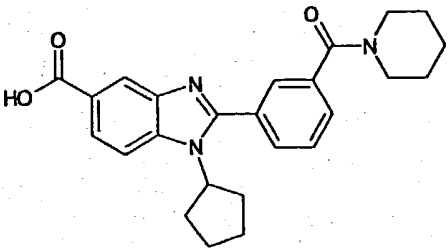
Ex. No.	Formula	MS
1289		509 (M+H)
1290		455 (M+H)
1291		494 (M+H)
1292		418 (M+H)



Table 125

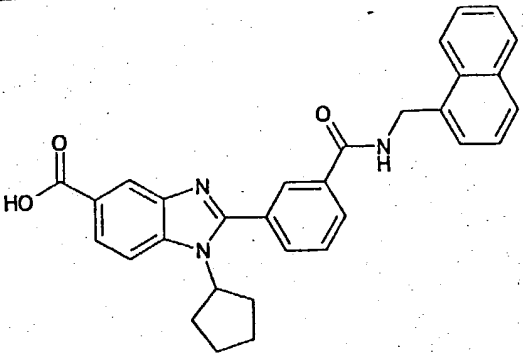
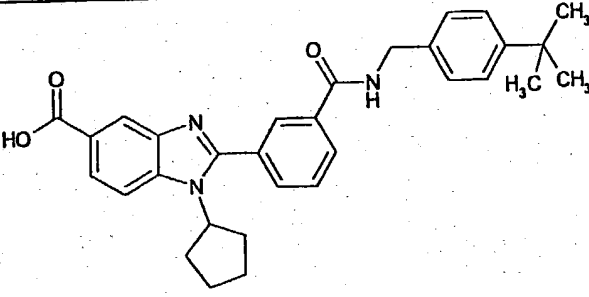
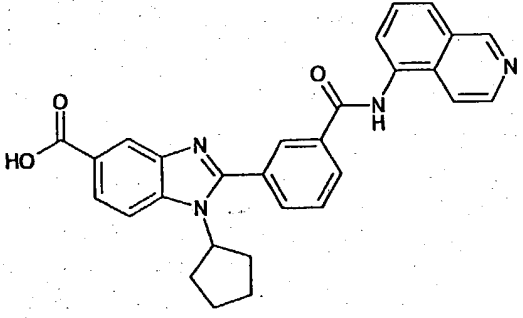
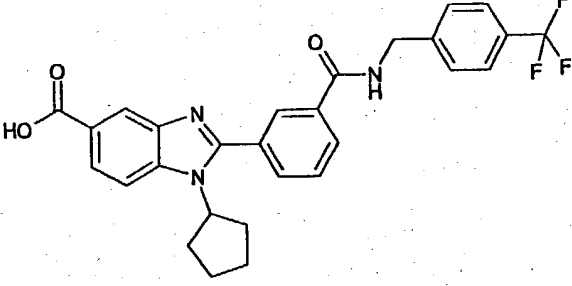
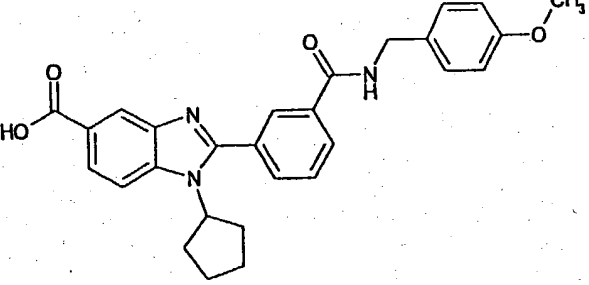
Ex. No.	Formula	MS
1293		490 (M+H)
1294		496 (M+H)
1295		477 (M+H)
1296		508 (M+H)
1297		470 (M+H)

Table 126

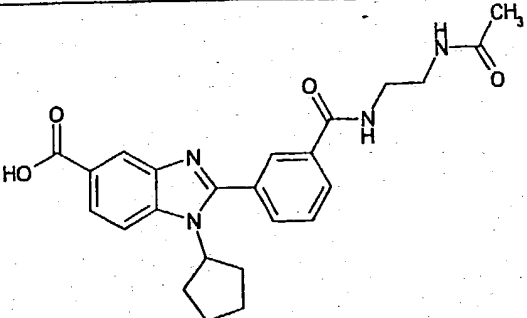
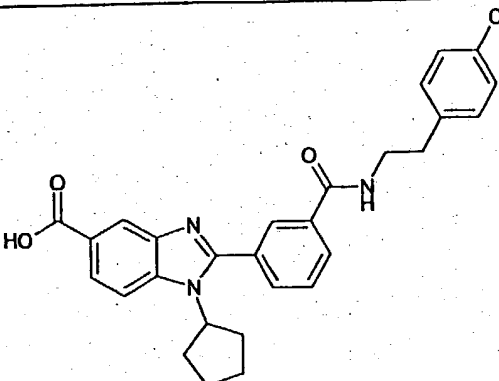
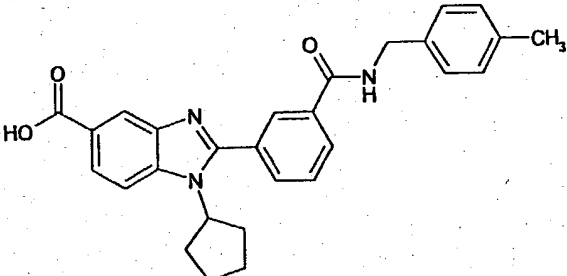
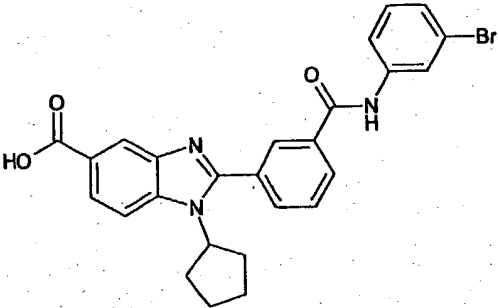
Ex. No.	Formula	MS
1298	 <chem>CC(=O)NCCC(=O)c1ccc(cc1)c2nc3cc(ccc3n2C4CCCC4)C(=O)O</chem>	435 (M+H)
1299	 <chem>Clc1ccc(cc1)CCNC(=O)c2ccc(cc2)c3nc4cc(ccc4n3C5CCCC5)C(=O)O</chem>	488 (M+H)
1300	 <chem>Cc1ccc(cc1)CCNC(=O)c2ccc(cc2)c3nc4cc(ccc4n3C5CCCC5)C(=O)O</chem>	454 (M+H)
1301	 <chem>BrC1=CC=C(C=C1)CCNC(=O)c2ccc(cc2)c3nc4cc(ccc4n3C5CCCC5)C(=O)O</chem>	504 (M+H)

Table 127

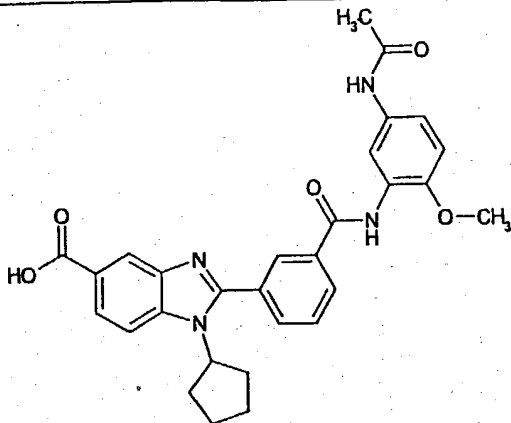
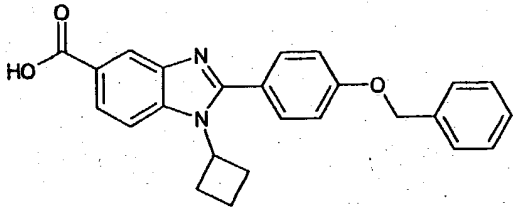
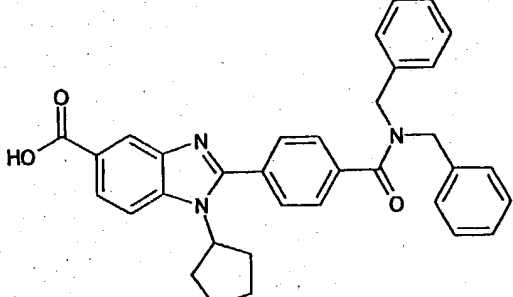
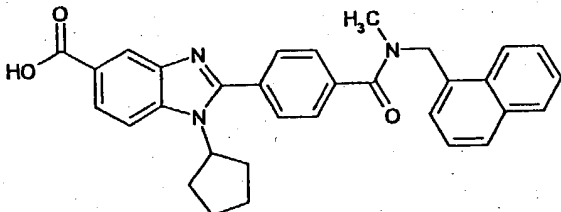
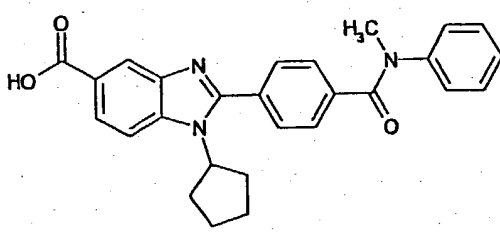
Ex. No.	Formula	MS
1302		513 (M+H)
1303		399 (M+H)
1304		530 (M+H)
1305		504 (M+H)
1306		440 (M+H)

Table 128

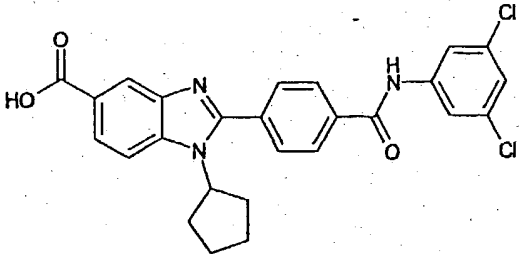
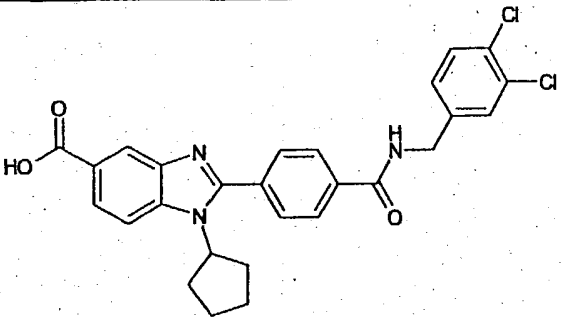
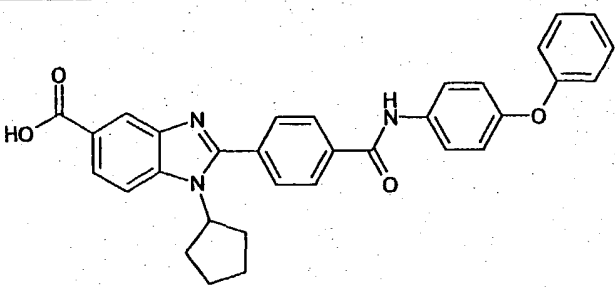
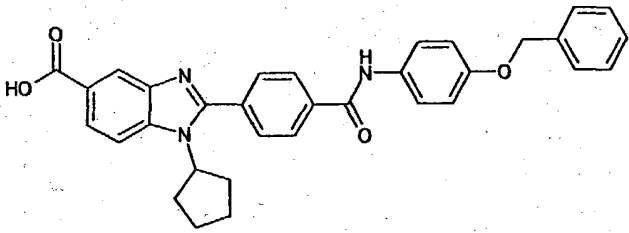
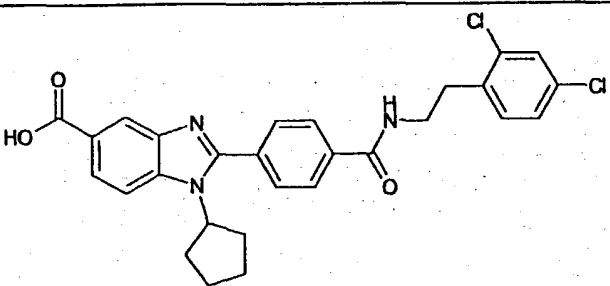
Ex. No.	Formula	MS
1307		494 (M+H)
1308		508 (M+H)
1309		518 (M+H)
1310		532 (M+H)
1311		522 (M+H)

Table 129

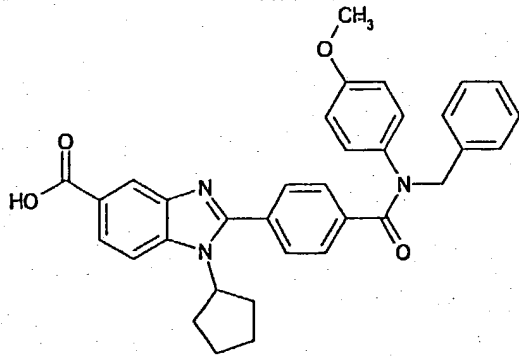
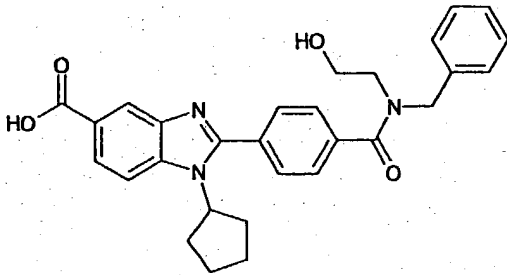
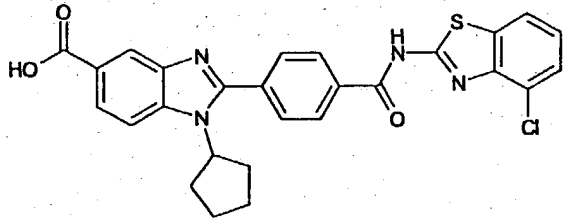
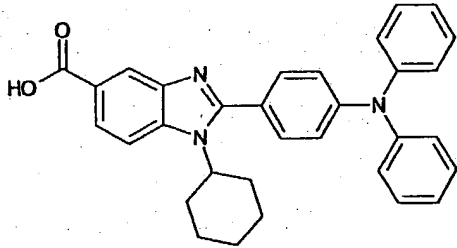
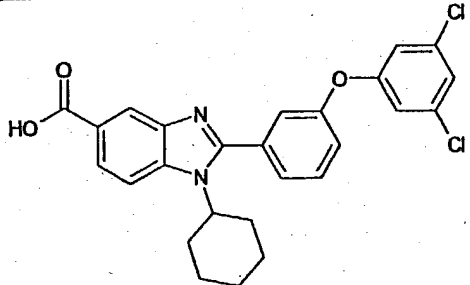
Ex. No.	Formula	MS
1312		546 (M+H)
1313		484 (M+H)
1314		517 (M+H)
1315		488 (M+H)
1316		481 (M+H)

Table 130

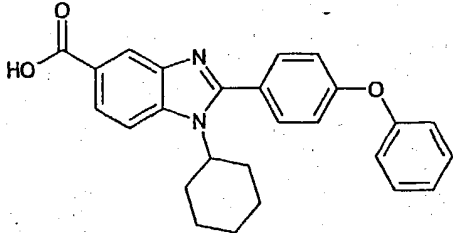
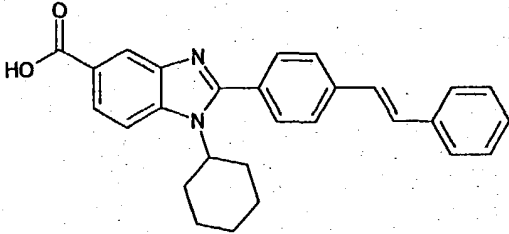
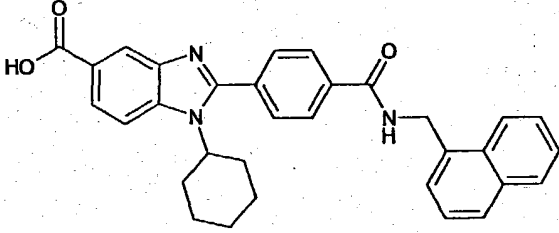
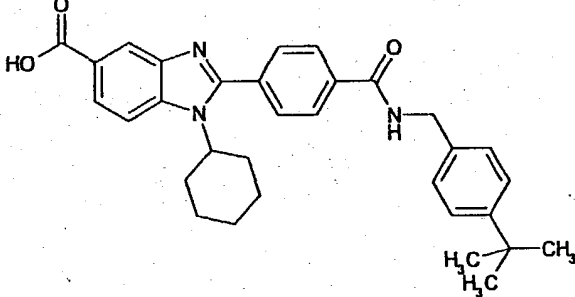
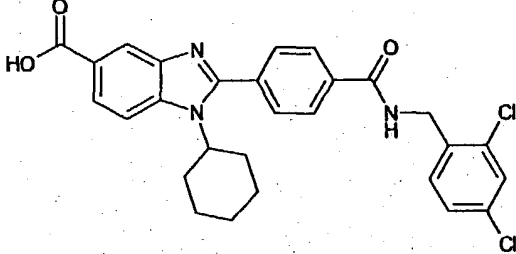
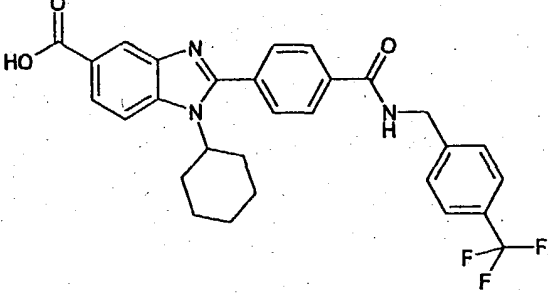
Ex. No.	Formula	MS
1317		413 (M+H)
1318		423 (M+H)
1319		504 (M+H)
1320		510 (M+H)
1321		522 (M+H)
1322		522 (M+H)

Table 131

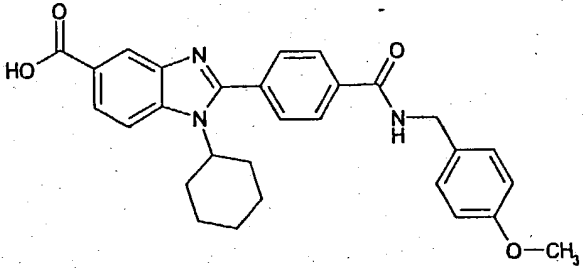
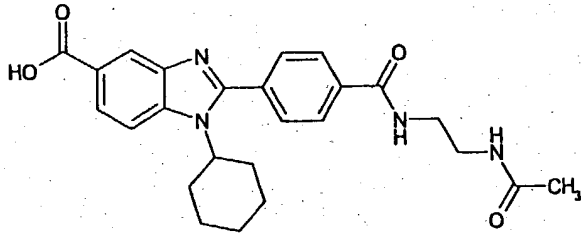
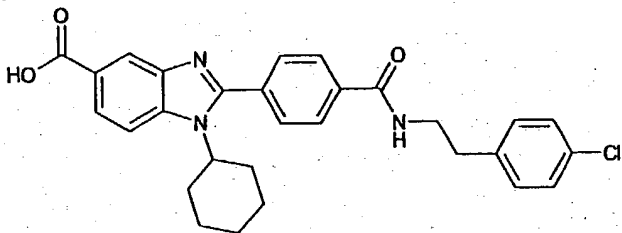
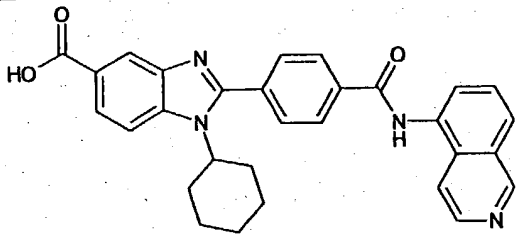
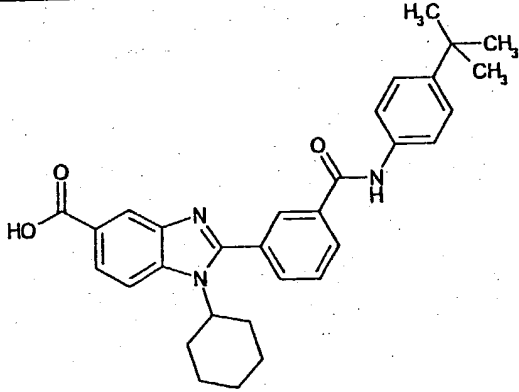
Ex. No.	Formula	MS
1323		484 (M+H)
1324		449 (M+H)
1325		502 (M+H)
1326		491 (M+H)
1327		496 (M+H)

Table 132

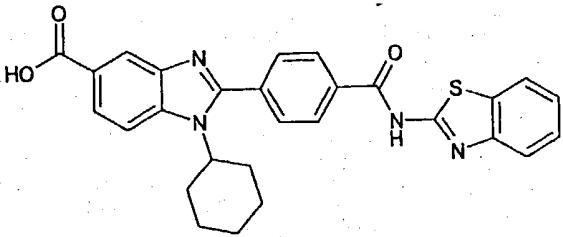
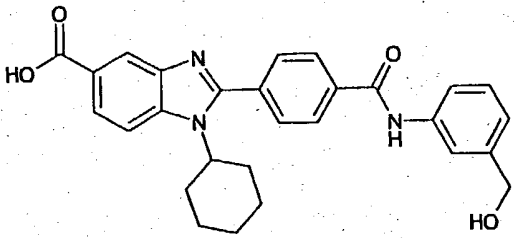
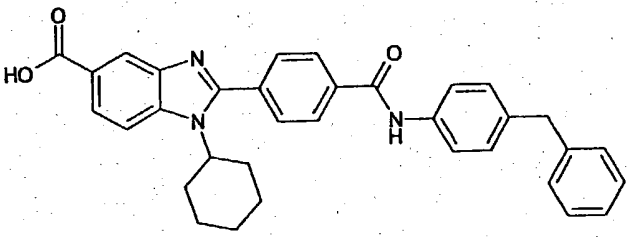
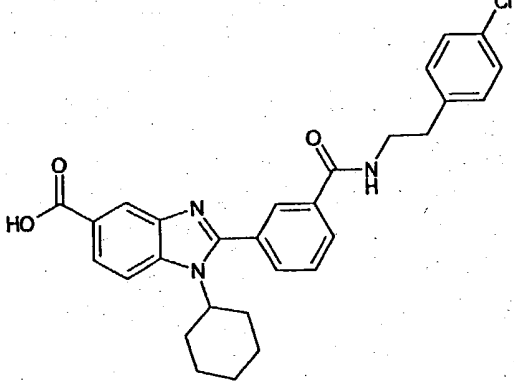
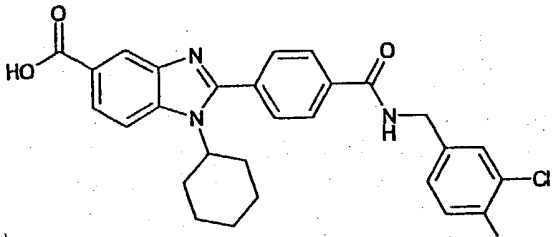
Ex. No.	Formula	MS
1328		497 (M+H)
1329		470 (M+H)
1330		530 (M+H)
1331		502 (M+H)
1332		522 (M+H)



Table 133

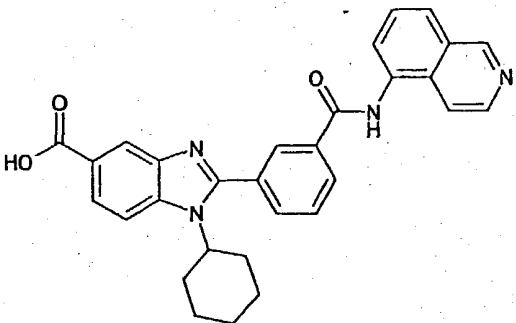
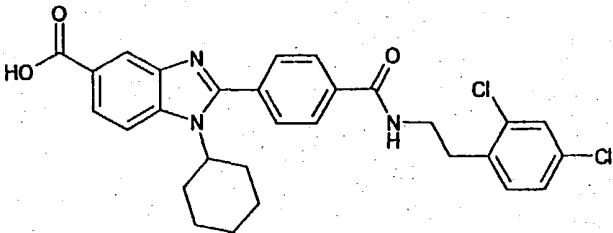
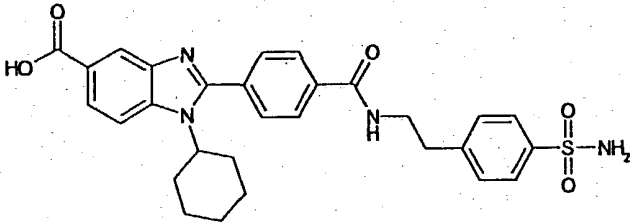
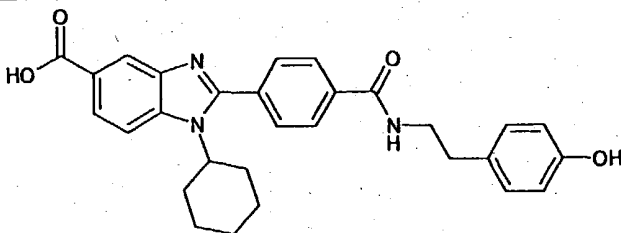
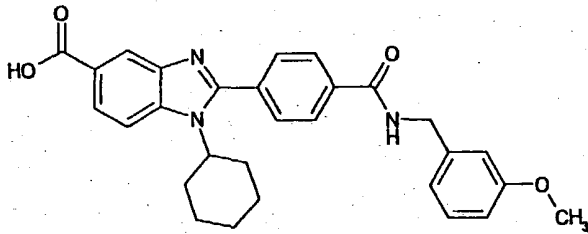
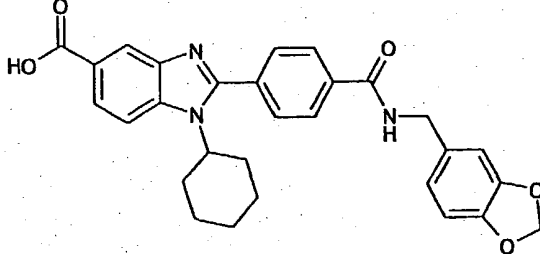
Ex. No.	Formula	MS
1333		491 (M+H)
1334		536 (M+H)
1335		547 (M+H)
1336		484 (M+H)
1337		484 (M+H)
1338		498 (M+H)

Table 134

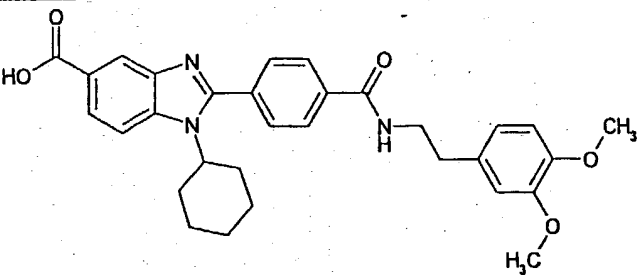
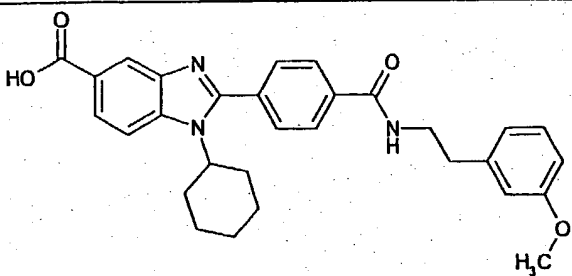
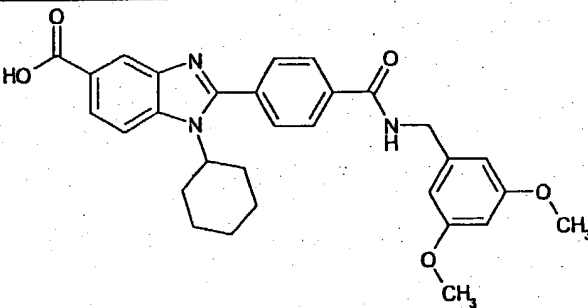
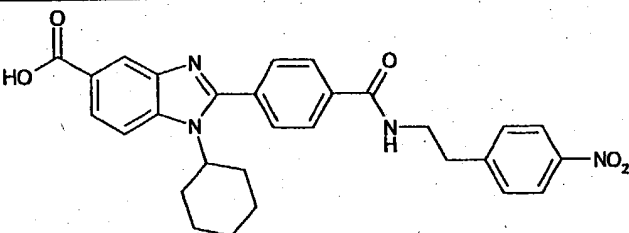
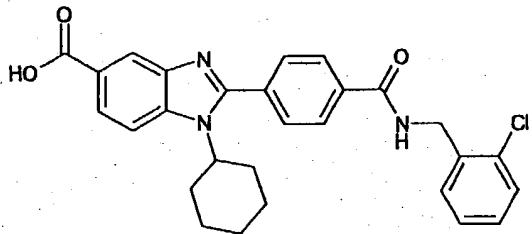
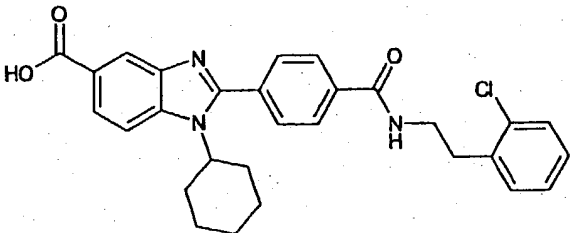
Ex. No.	Formula	MS
1339		528 (M+H)
1340		498 (M+H)
1341		514 (M+H)
1342		513 (M+H)
1343		488 (M+H)
1344		502 (M+H)

Table 135

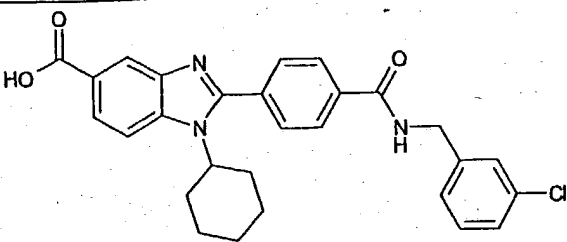
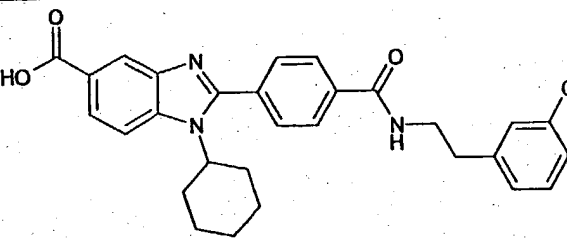
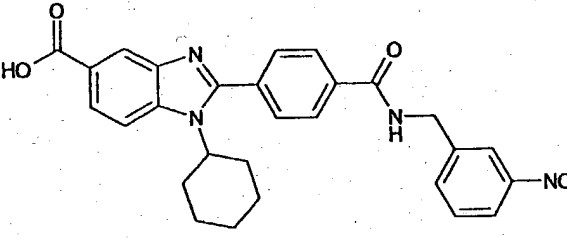
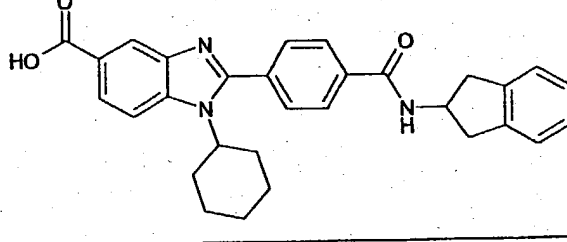
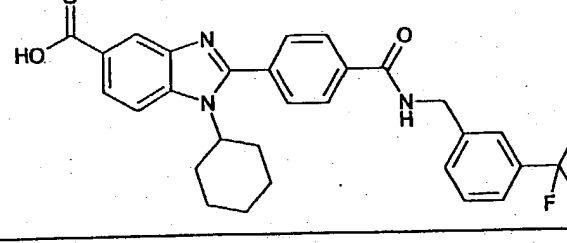
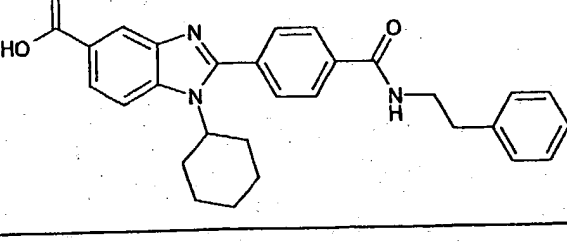
Ex. No.	Formula	MS
1345		488 (M+H)
1346		502 (M+H)
1347		499 (M+H)
1348		480 (M+H)
1349		522 (M+H)
1350		546 (M+H)

Table 136

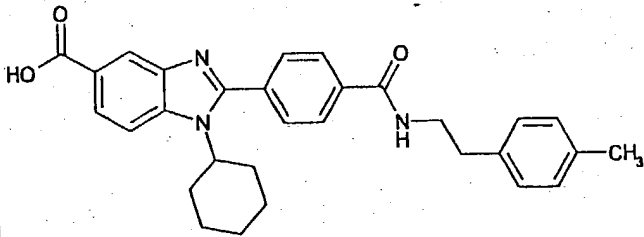
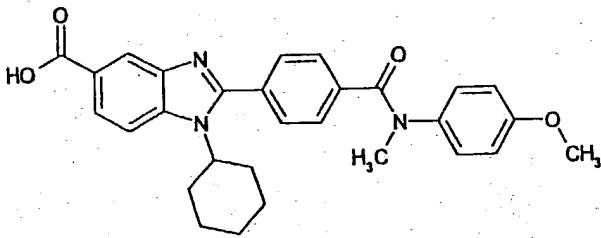
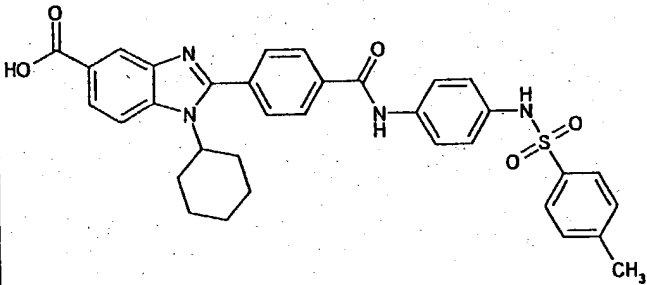
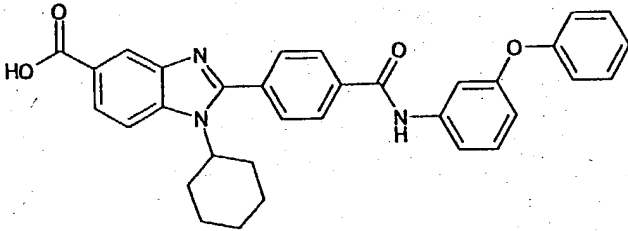
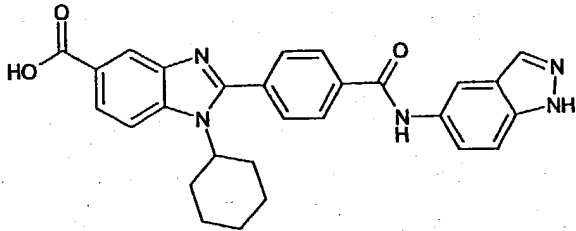
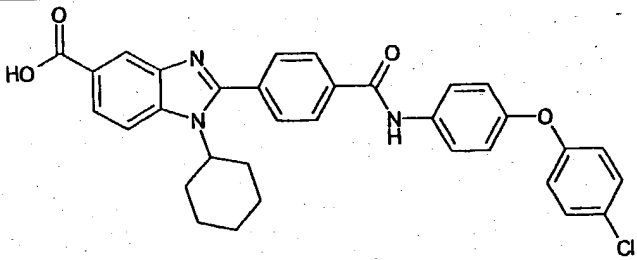
Ex. No.	Formula	MS
1351		482 (M+H)
1352		484 (M+H)
1353		609 (M+H)
1354		532 (M+H)
1355		480 (M+H)
1356		566 (M+H)

Table 137

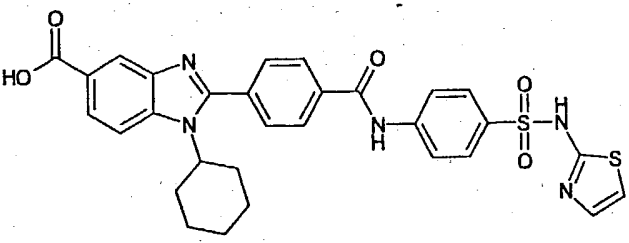
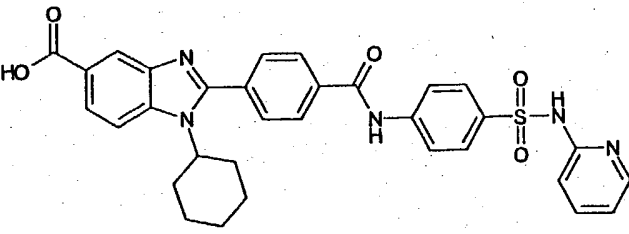
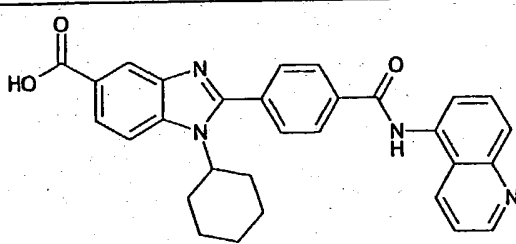
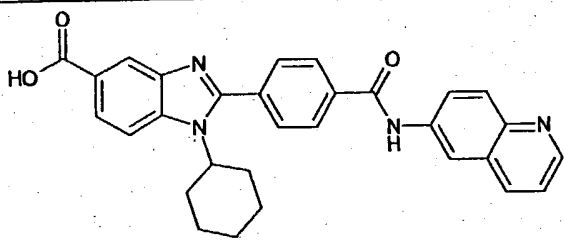
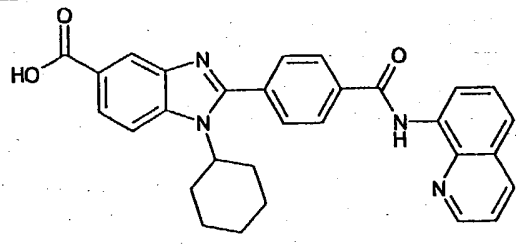
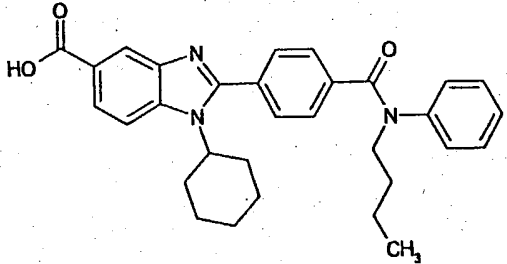
Ex. No.	Formula	MS
1357		602 (M+H)
1358		596 (M+H)
1359		491 (M+H)
1360		491 (M+H)
1361		491 (M+H)
1362		496 (M+H)

Table 138

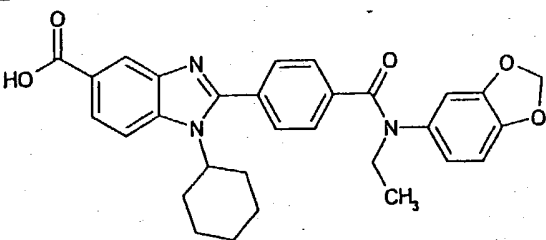
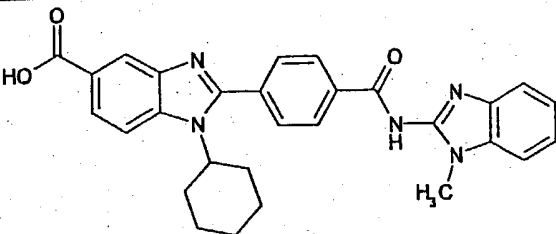
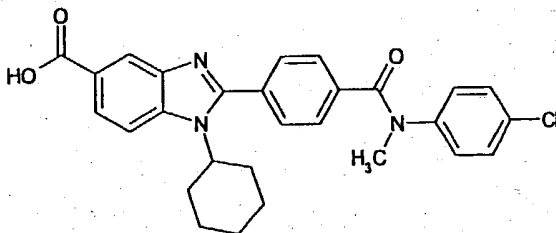
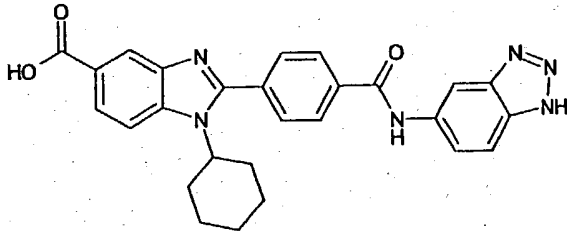
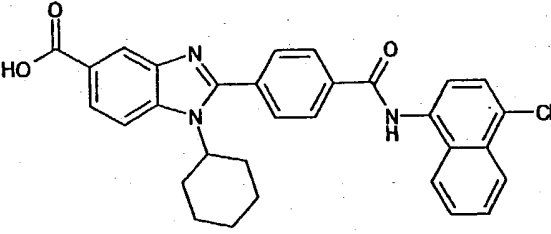
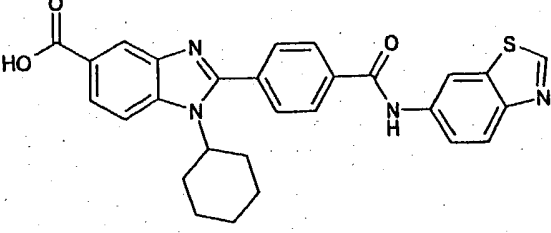
Ex. No.	Formula	MS
1363		512 (M+H)
1364		494 (M+H)
1365		488 (M+H)
1366		481 (M+H)
1367		524 (M+H)
1368		497 (M+H)

Table 139

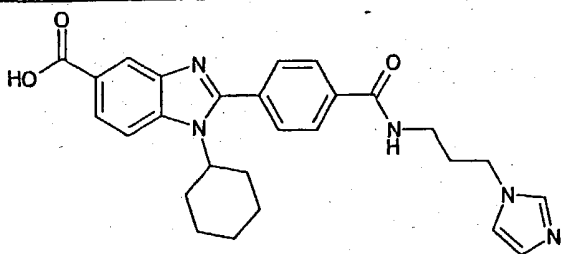
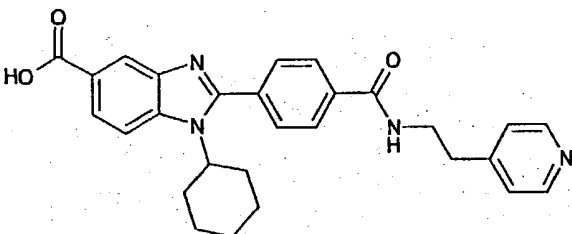
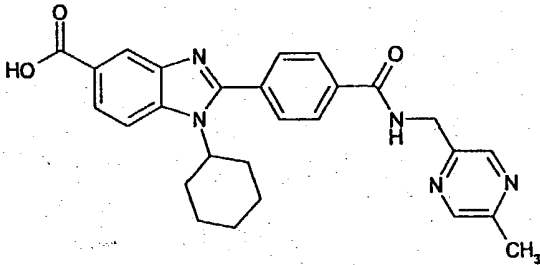
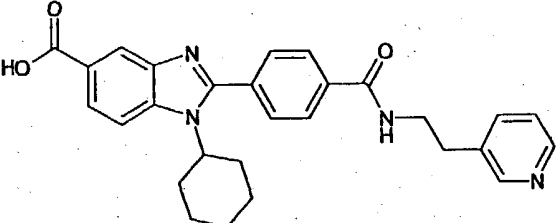
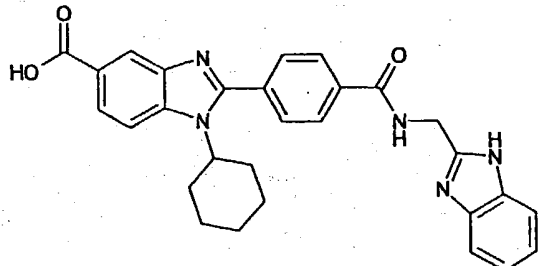
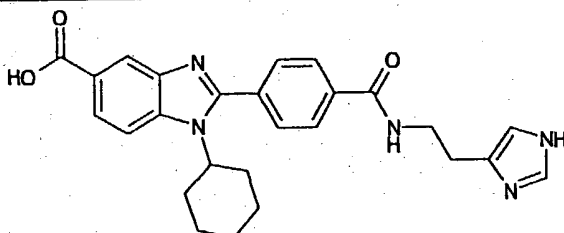
Ex. No.	Formula	MS
1369		472 (M+H)
1370		469 (M+H)
1371		470 (M+H)
1372		469 (M+H)
1373		494 (M+H)
1374		458 (M+H)

Table 140

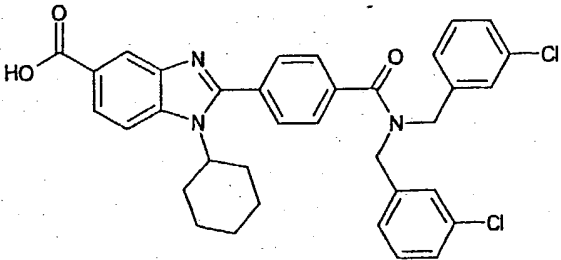
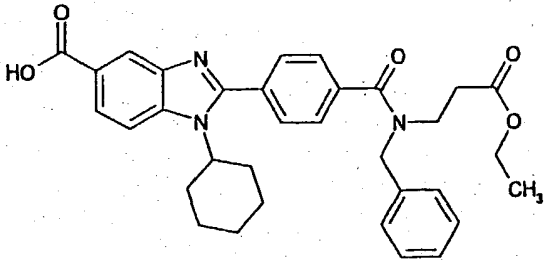
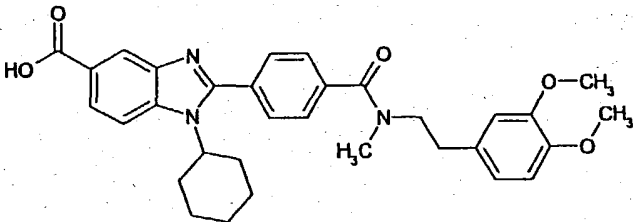
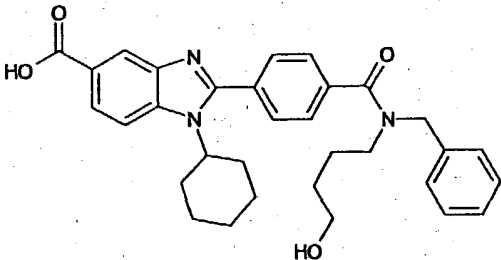
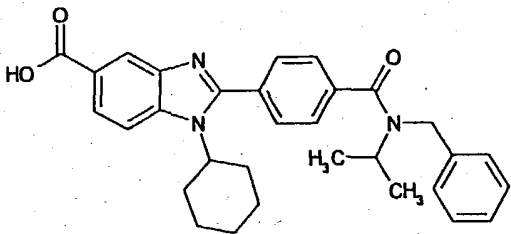
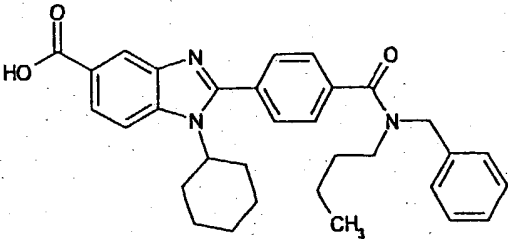
Ex. No.	Formula	MS
1375		612 (M+H)
1376		554 (M+H)
1377		542 (M+H)
1378		526 (M+H)
1379		496 (M+H)
1380		510 (M+H)



Table 141

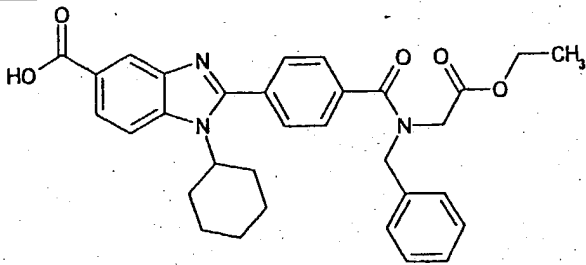
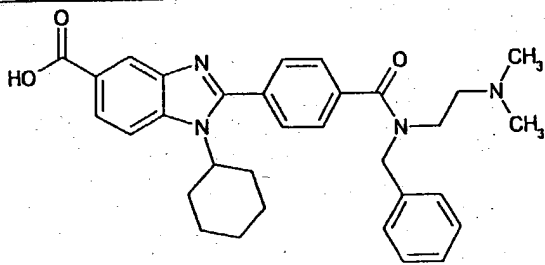
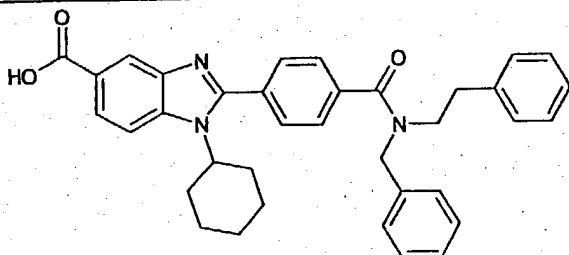
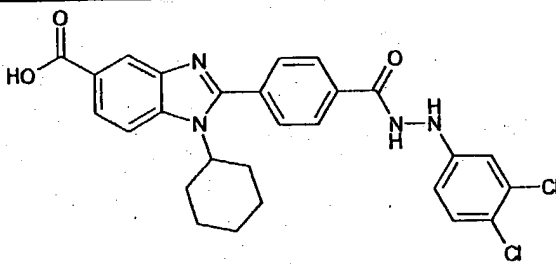
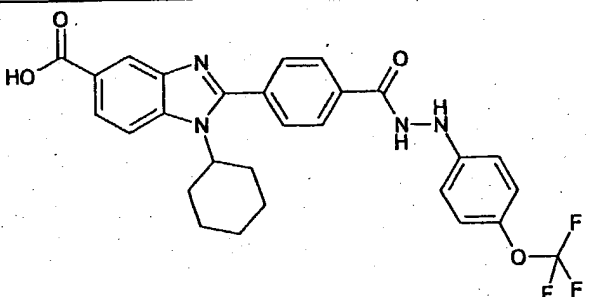
Ex. No.	Formula	MS
1381		540 (M+H)
1382		525 (M+H)
1383		558 (M+H)
1384		523 (M+H)
1385		539 (M+H)

Table 142

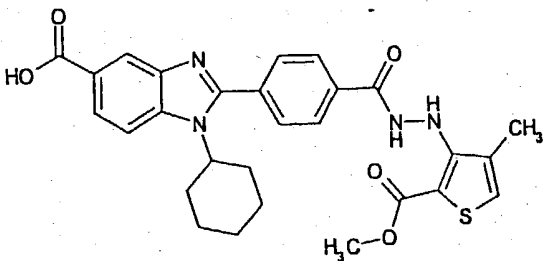
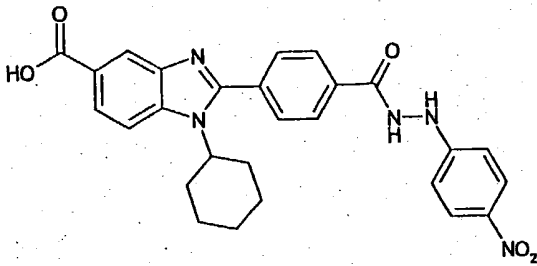
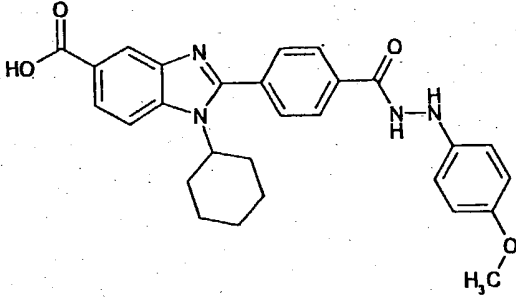
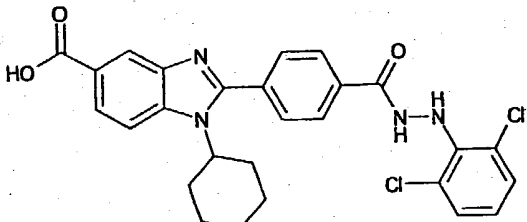
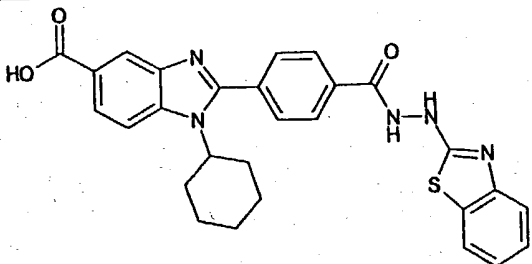
Ex. No.	Formula	MS
1386		533 (M+H)
1387		500 (M+H)
1388		485 (M+H)
1389		523 (M+H)
1390		512 (M+H)

Table 143

Ex. No.	Formula	MS
1391		540 (M+H)
1392		527 (M+H)
1393		525 (M+H)
1394		507 (M+H)
1395		491 (M+H)
1396		506 (M+H)

Table 144

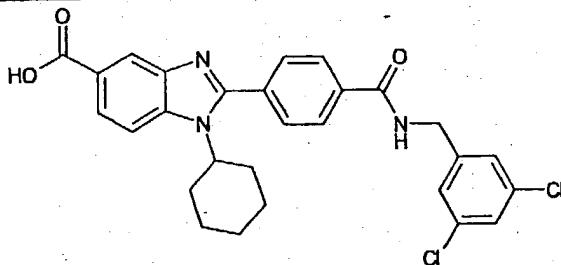
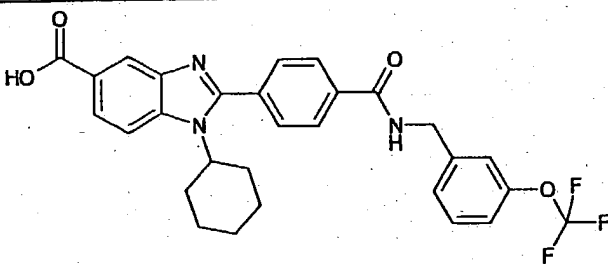
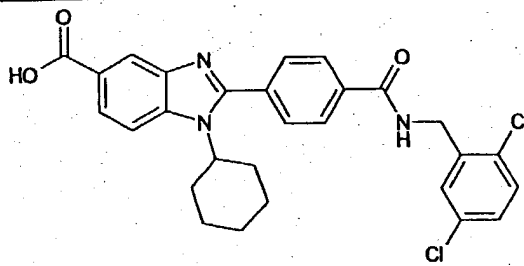
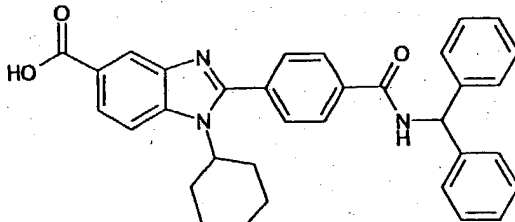
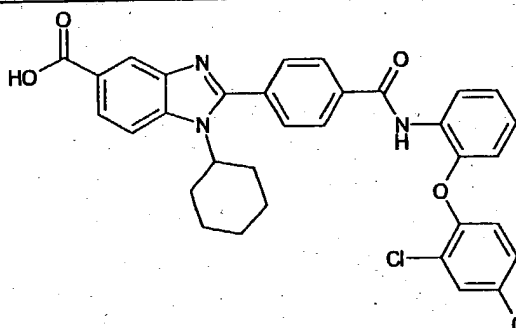
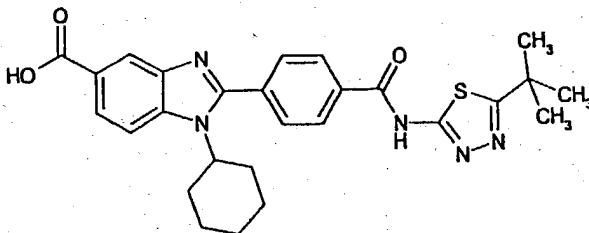
Ex. No.	Formula	MS
1397		522 (M+H)
1398		538 (M+H)
1399		522 (M+H)
1400		530 (M+H)
1401		600 (M+H)
1402		504 (M+H)

Table 145

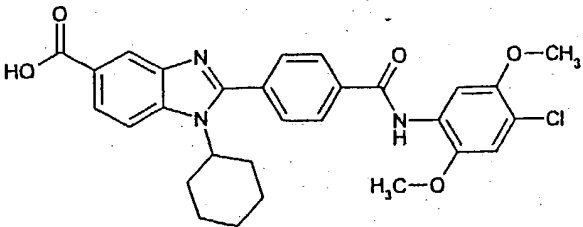
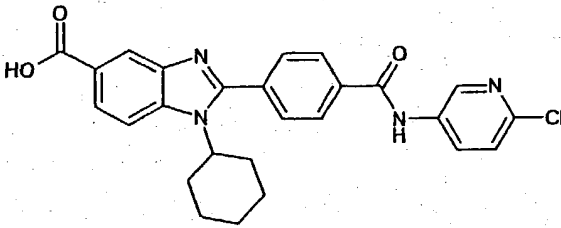
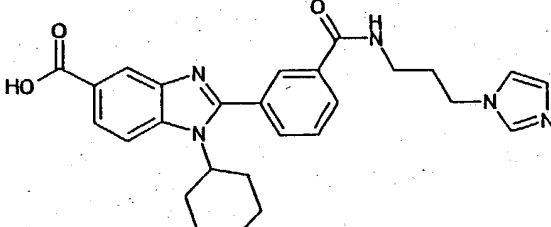
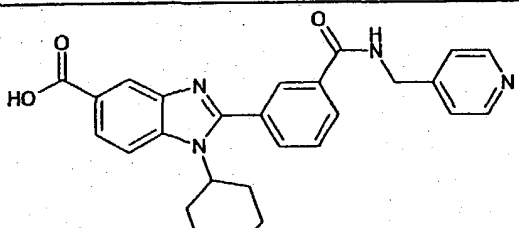
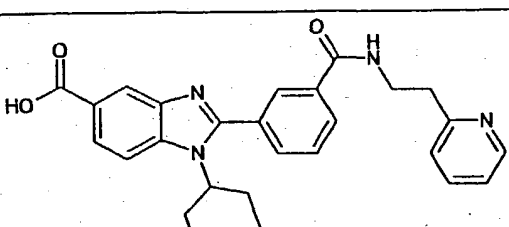
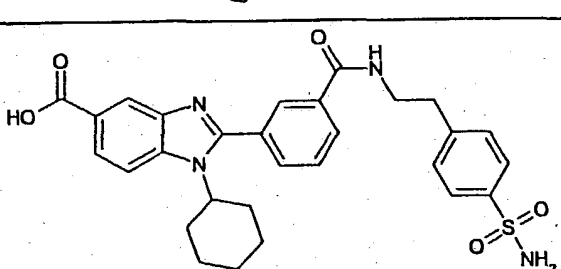
Ex. No.	Formula	MS
1403		534 (M+H)
1404		475 (M+H)
1405		472 (M+H)
1406		455 (M+H)
1407		469 (M+H)
1408		547 (M+H)

Table 146

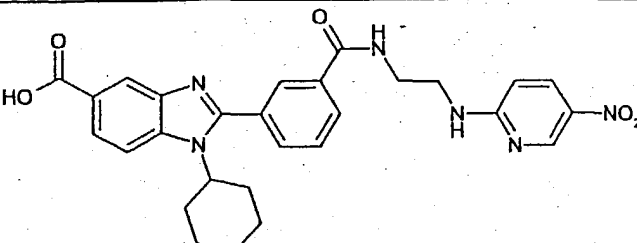
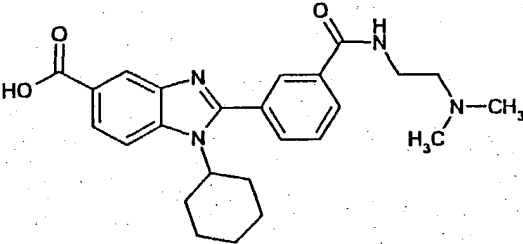
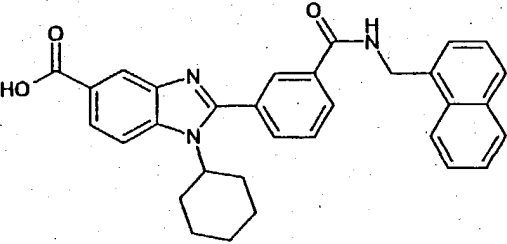
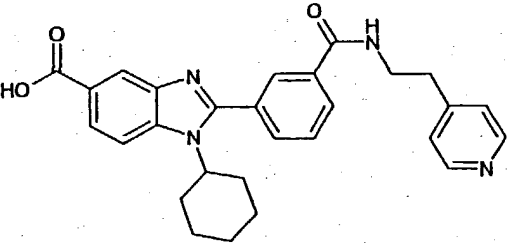
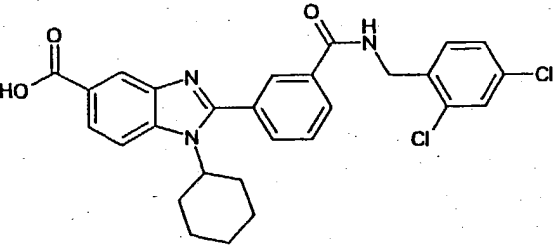
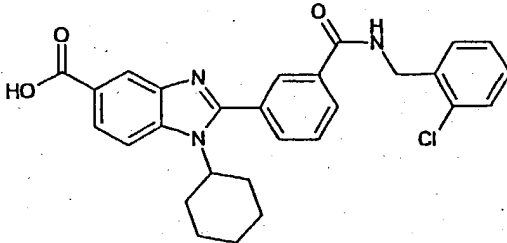
Ex. No.	Formula	MS
1409		529 (M+H)
1410		435 (M+H)
1411		504 (M+H)
1412		469 (M+H)
1413		522 (M+H)
1414		488 (M+H)

Table 147

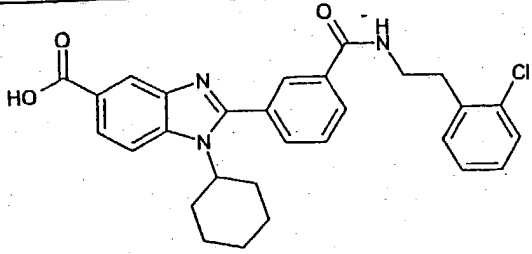
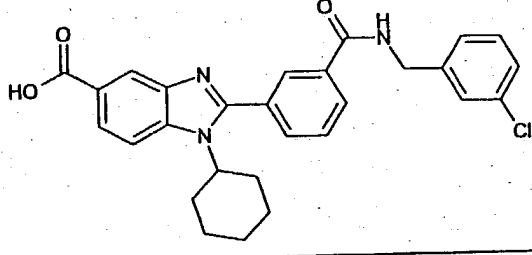
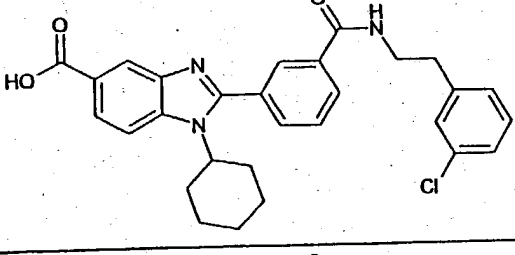
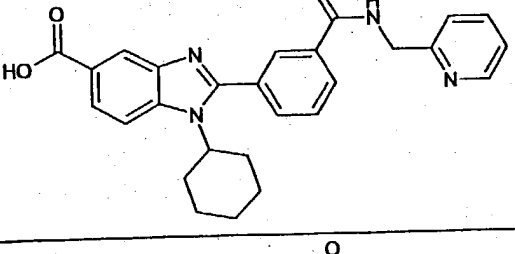
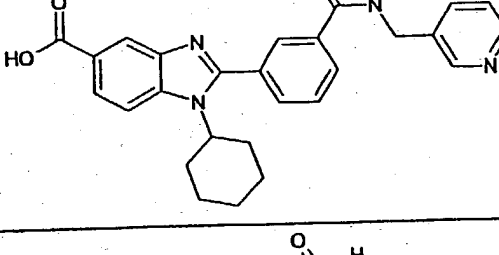
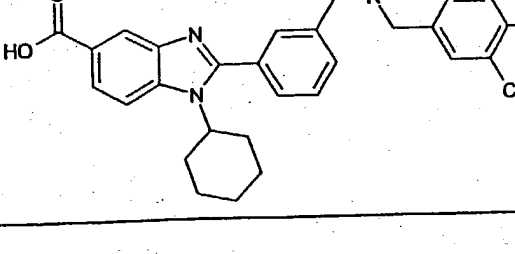
Ex. No.	Formula	MS
1415		502 (M+H)
1416		488 (M+H)
1417		502 (M+H)
1418		455 (M+H)
1419		455 (M+H)
1420		522 (M+H)

Table 148

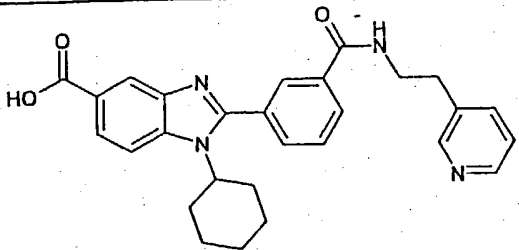
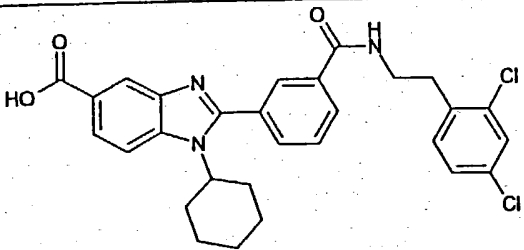
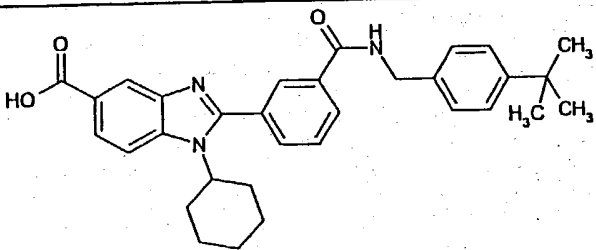
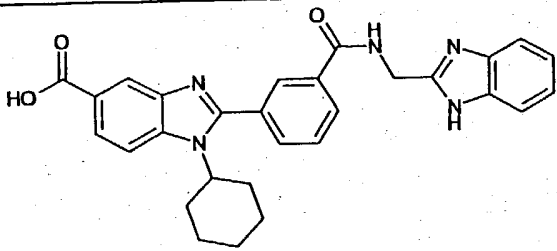
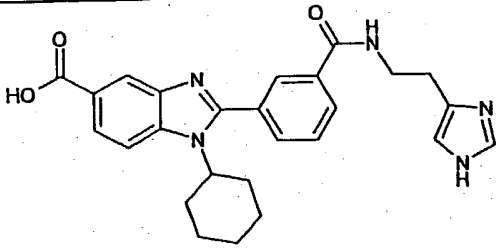
Ex. No.	Formula	MS
1421		469 (M+H)
1422		536 (M+H)
1423		510 (M+H)
1424		494 (M+H)
1425		458 (M+H)



Table 149

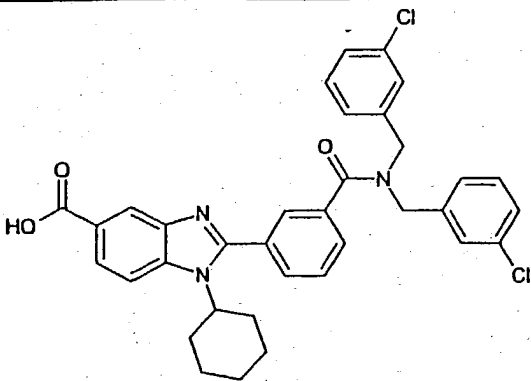
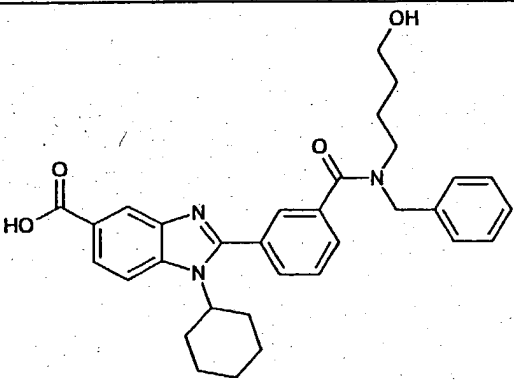
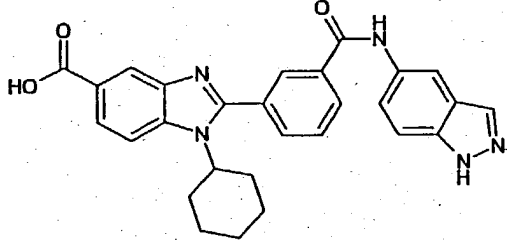
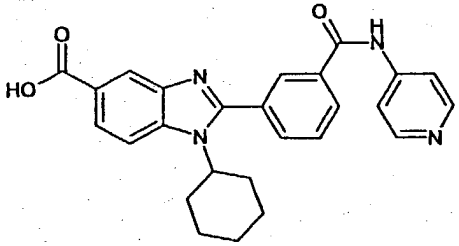
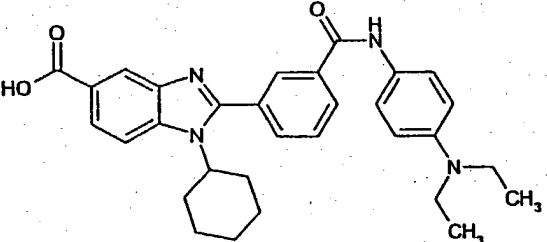
Ex. No.	Formula	MS
1426		612 (M+H)
1427		526 (M+H)
1428		480 (M+H)
1429		441 (M+H)
1430		511 (M+H)

Table 150

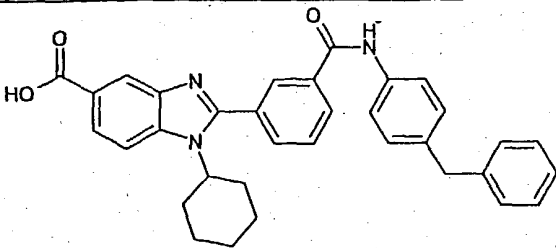
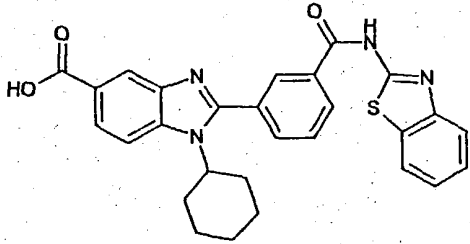
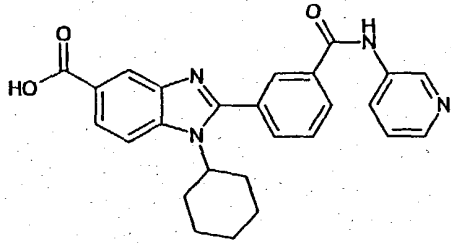
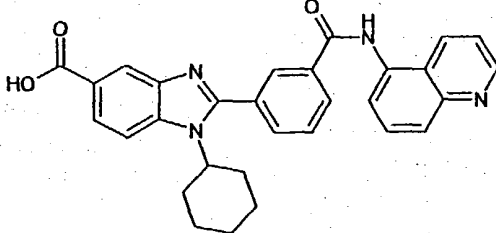
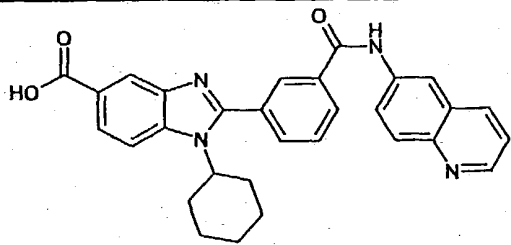
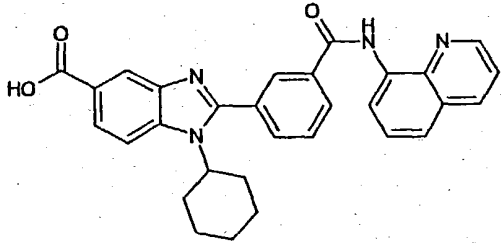
Ex. No.	Formula	MS
1431		530 (M+H)
1432		497 (M+H)
1433		441 (M+H)
1434		491 (M+H)
1435		491 (M+H)
1436		491 (M+H)

Table 151

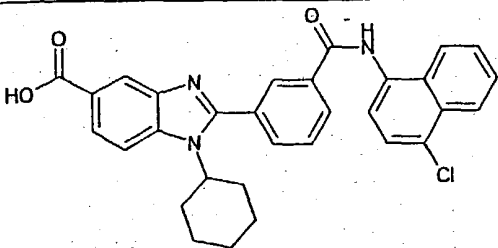
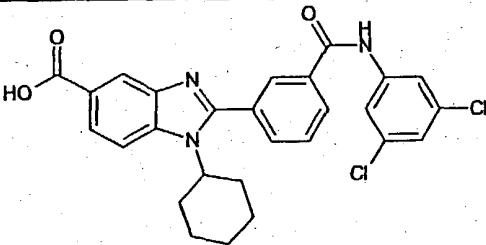
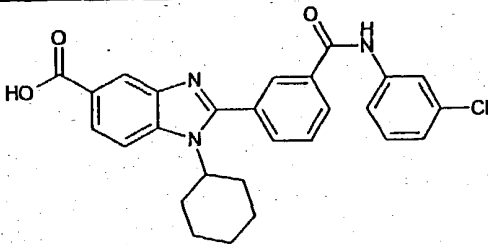
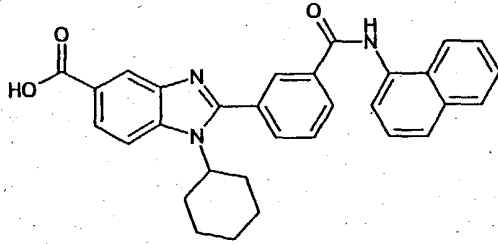
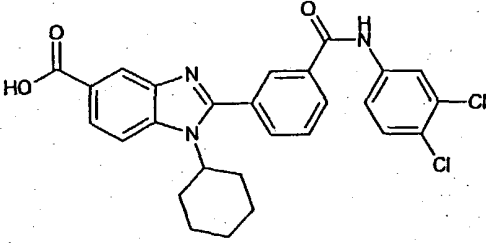
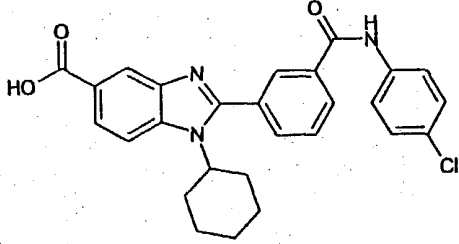
Ex. No.	Formula	MS
1437		524 (M+H)
1438		508 (M+H)
1439		474 (M+H)
1440		490 (M+H)
1441		508 (M+H)
1442		474 (M+H)

Table 152

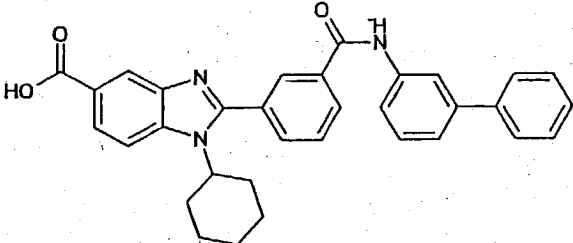
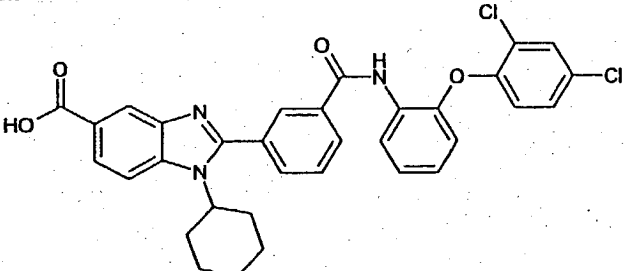
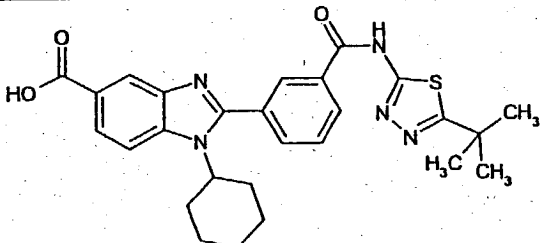
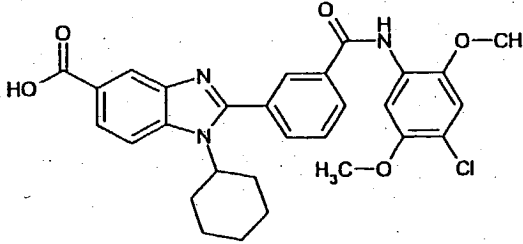
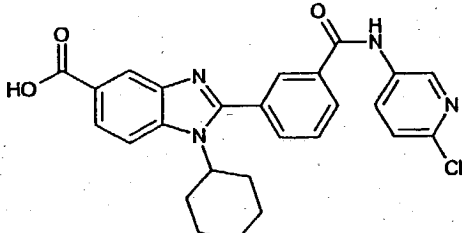
Ex. No.	Formula	MS
1443		516 (M+H)
1444		600 (M+H)
1445		504 (M+H)
1446		534 (M+H)
1447		475 (M+H)

Table 153

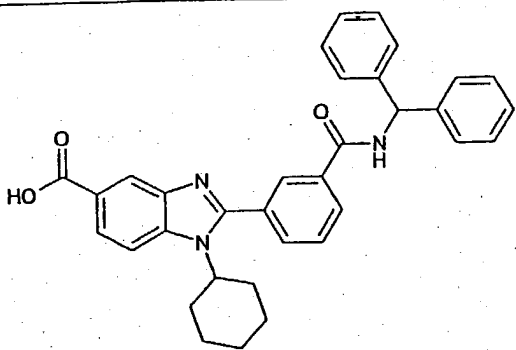
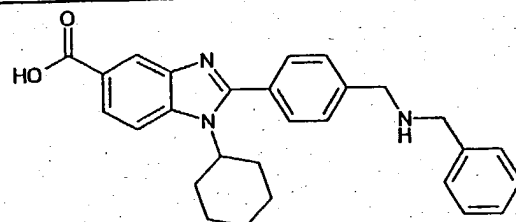
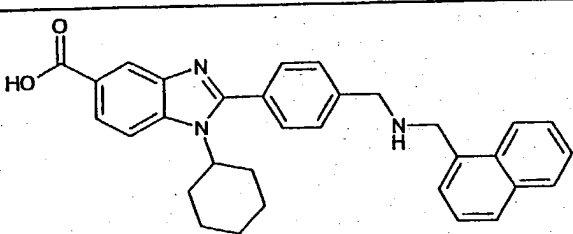
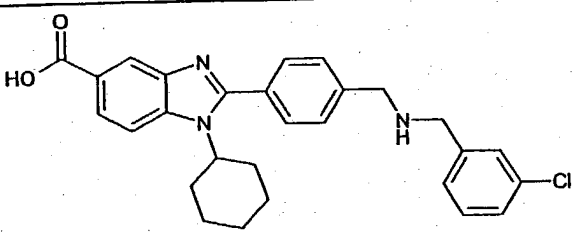
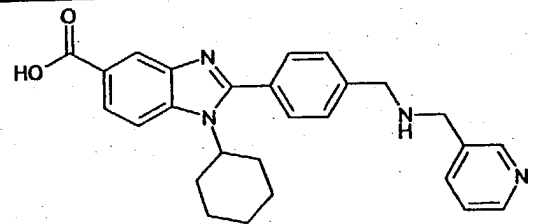
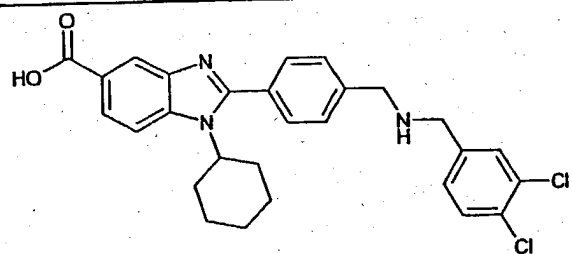
Ex. No.	Formula	MS
1448		530 (M+H)
1449		440 (M+H)
1450		490 (M+H)
1451		474 (M+H)
1452		441 (M+H)
1453		508 (M+H)

Table 154

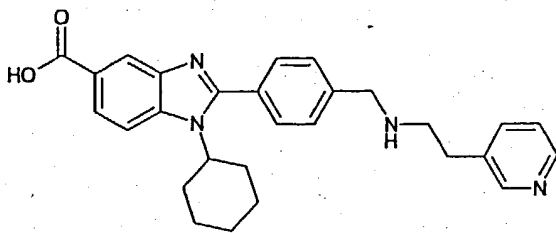
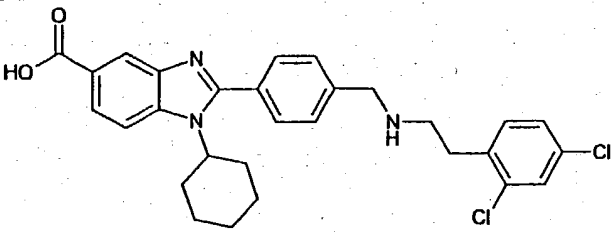
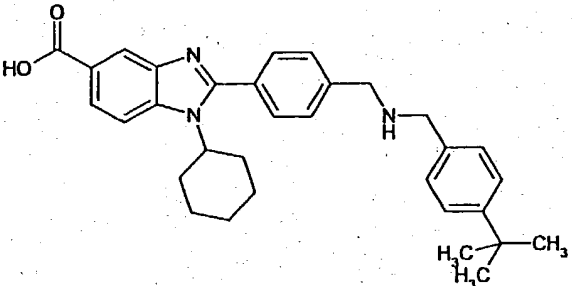
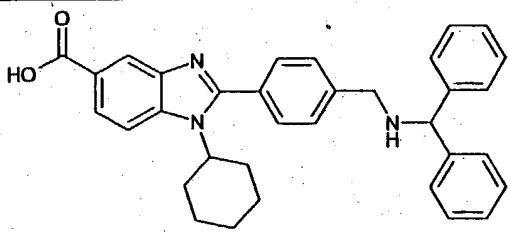
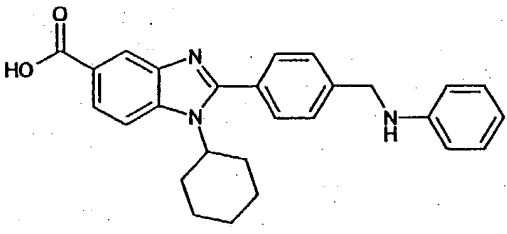
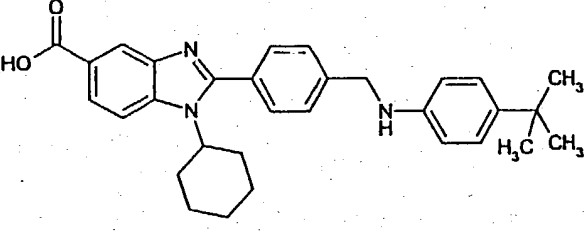
Ex. No.	Formula	MS
1454		455 (M+H)
1455		522 (M+H)
1456		496 (M+H)
1457		516 (M+H)
1458		426 (M+H)
1459		482 (M+H)

Table 155

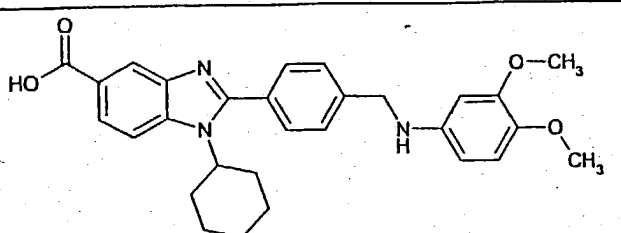
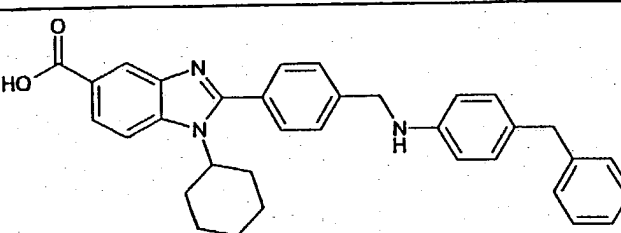
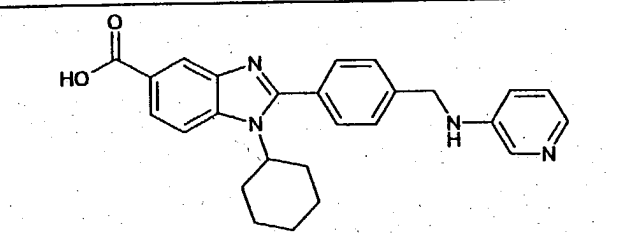
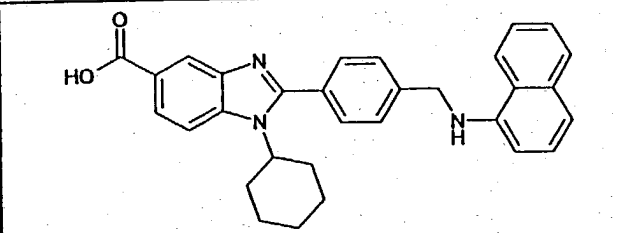
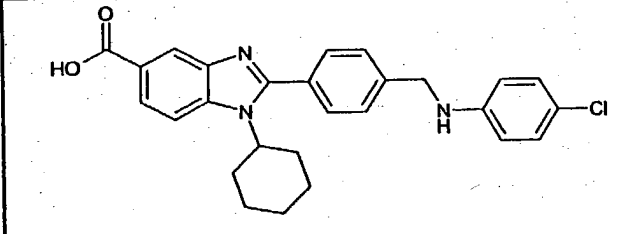
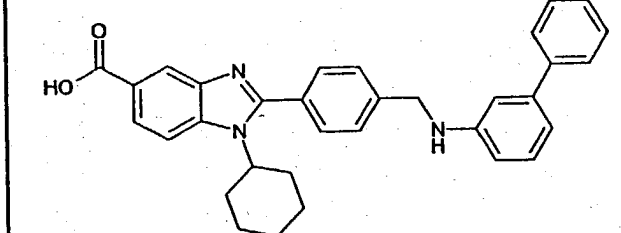
Ex. No.	Formula	MS
1460		486 (M+H)
1461		516 (M+H)
1462		427 (M+H)
1463		476 (M+H)
1464		460 (M+H)
1465		502 (M+H)

Table 156

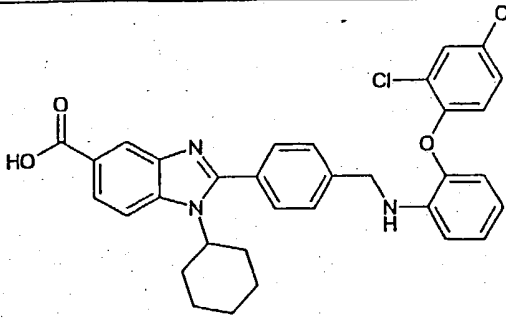
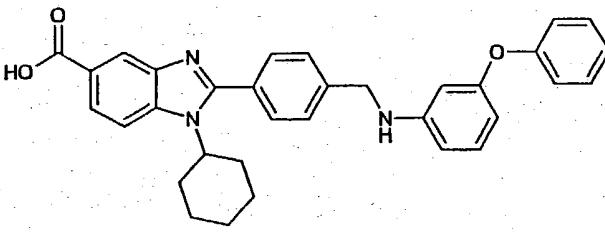
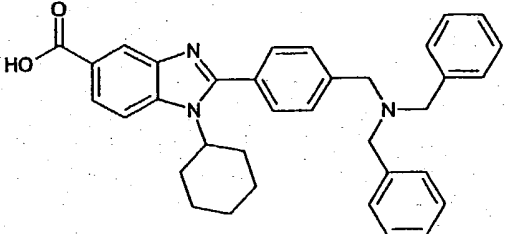
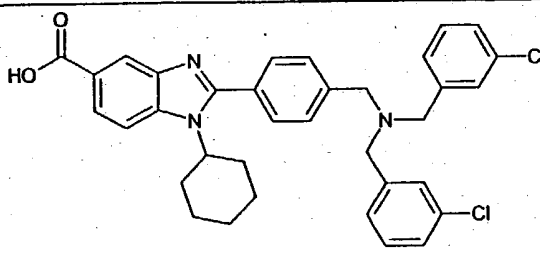
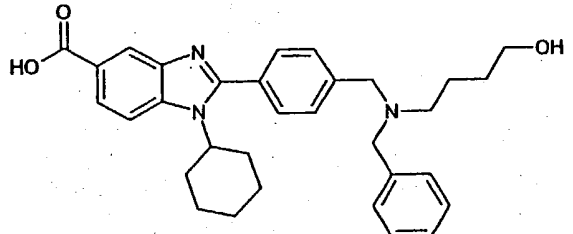
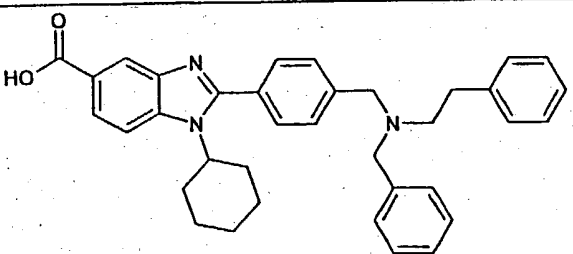
Ex. No.	Formula	MS
1466		586 (M+H)
1467		518 (M+H)
1468		530 (M+H)
1469		598 (M+H)
1470		512 (M+H)
1471		544 (M+H)



Table 157

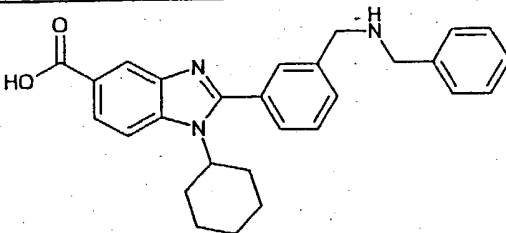
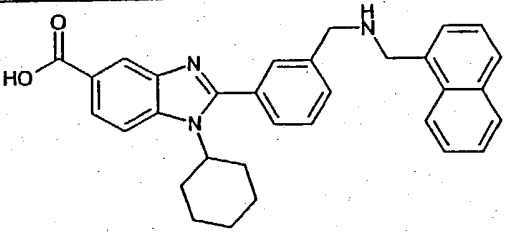
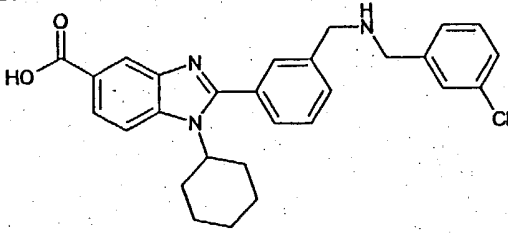
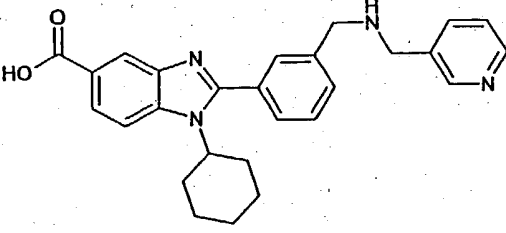
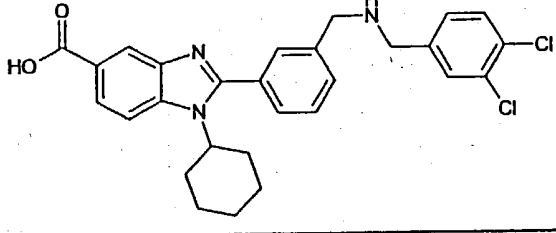
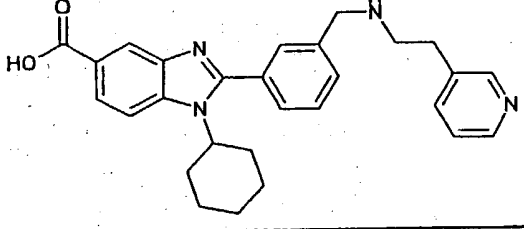
Ex. No.	Formula	MS
1472		440 (M+H)
1473		490 (M+H)
1474		474 (M+H)
1475		441 (M+H)
1476		508 (M+H)
1477		455 (M+H)

Table 158

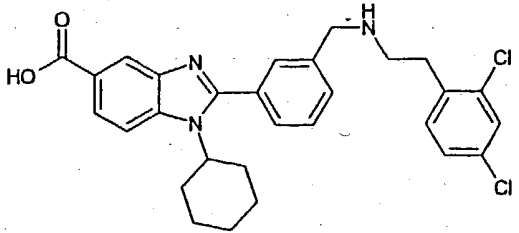
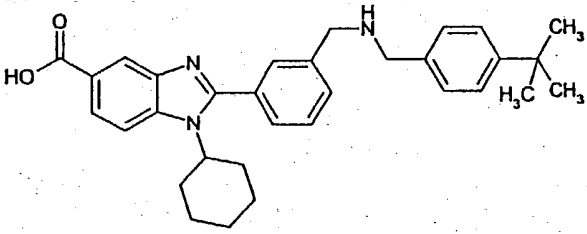
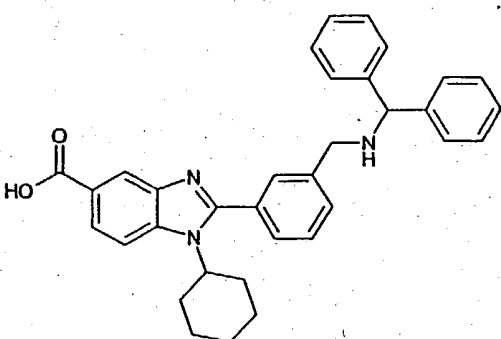
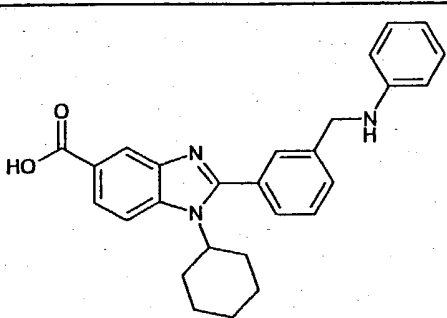
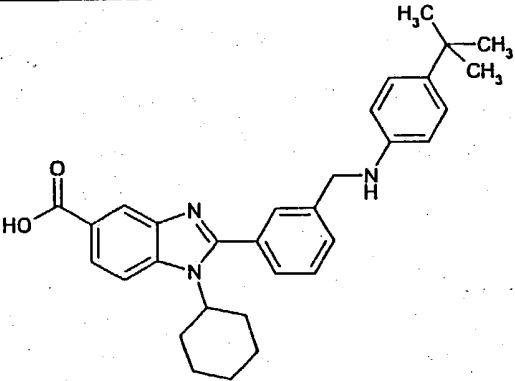
Ex. No.	Formula	MS
1478		522 (M+H)
1479		496 (M+H)
1480		516 (M+H)
1481		426 (M+H)
1482		482 (M+H)

Table 159

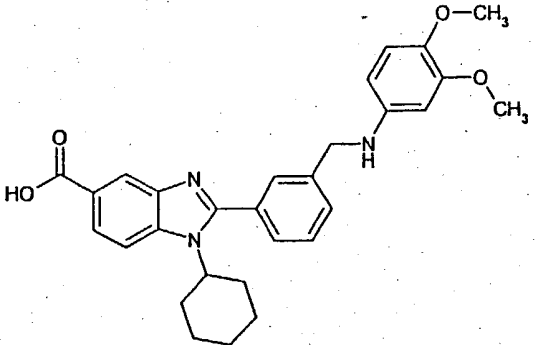
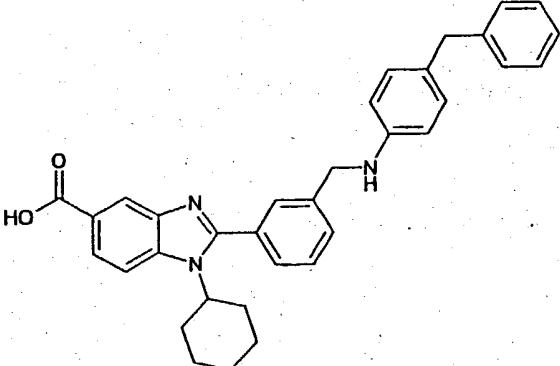
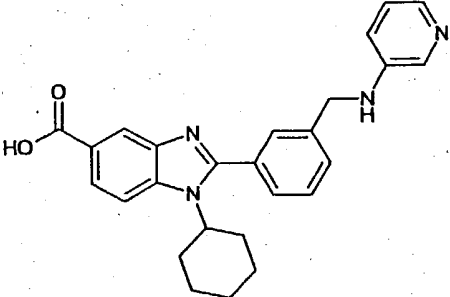
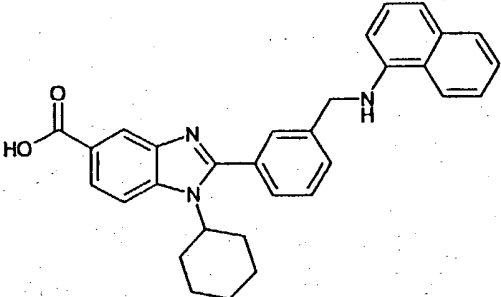
Ex. No.	Formula	MS
1483		486 (M+H)
1484		516 (M+H)
1485		427 (M+H)
1486		476 (M+H)

Table 160

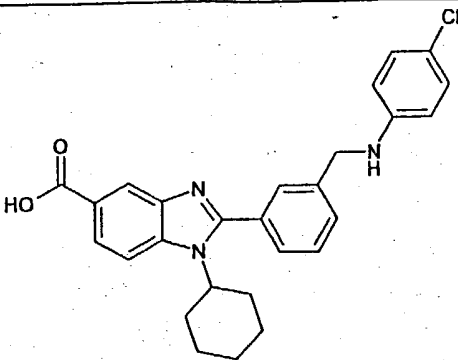
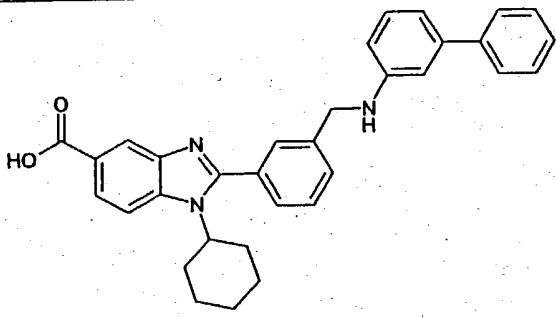
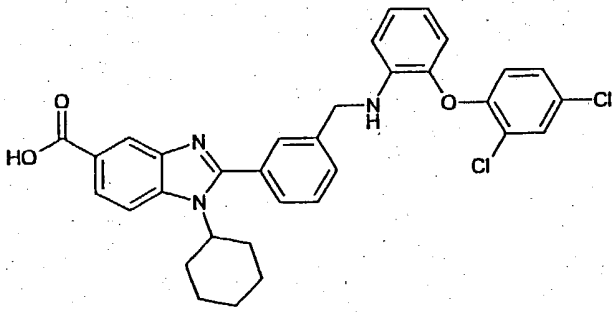
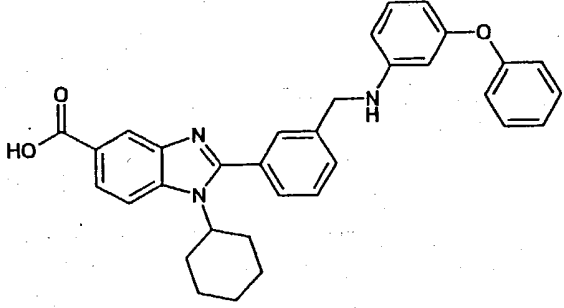
Ex. No.	Formula	MS
1487		460 (M+H)
1488		502 (M+H)
1489		586 (M+H)
1490		518 (M+H)

Table 161.

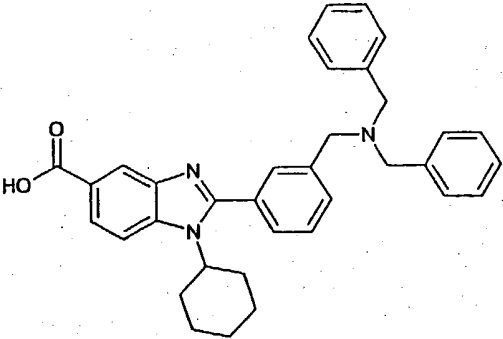
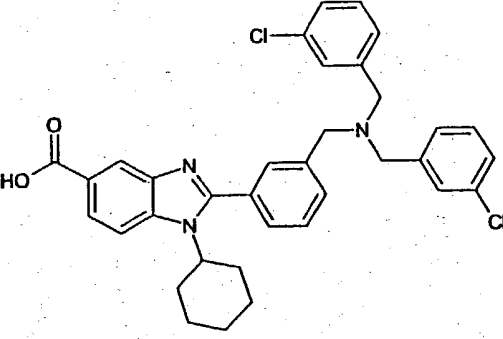
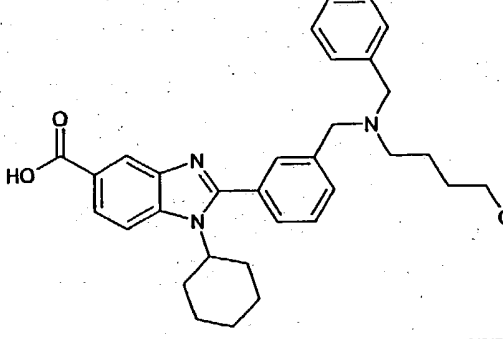
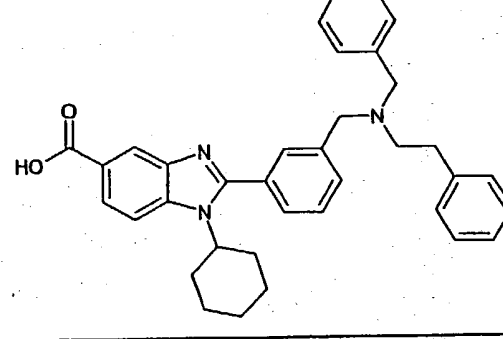
Ex. No.	Formula	MS
1491		530 (M+H)
1492		598 (M+H)
1493		512 (M+H)
1494		544 (M+H)

Table 162

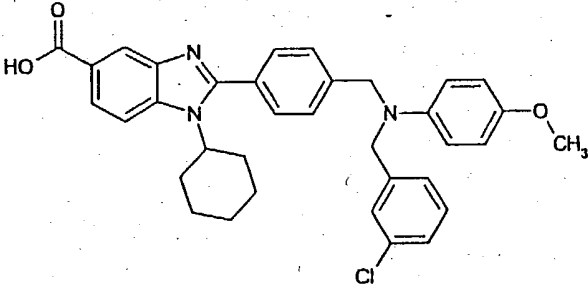
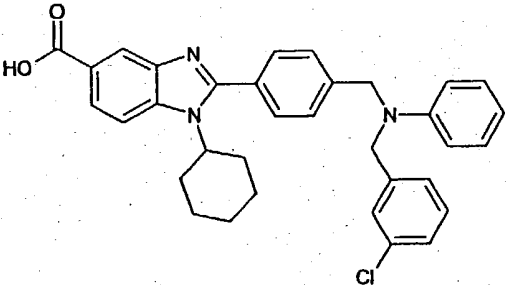
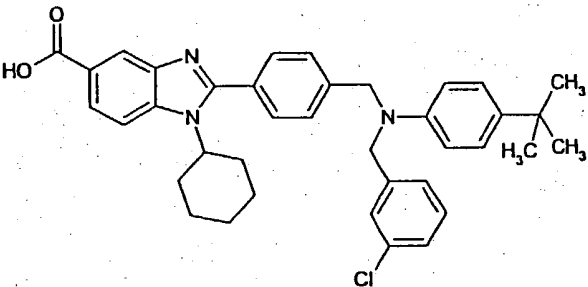
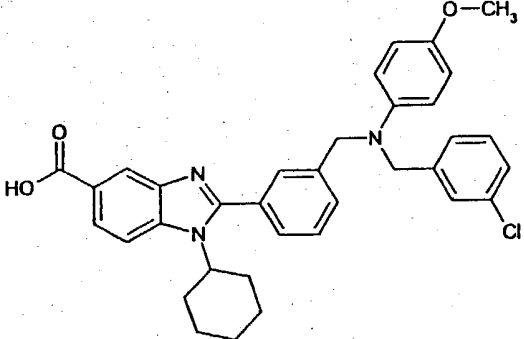
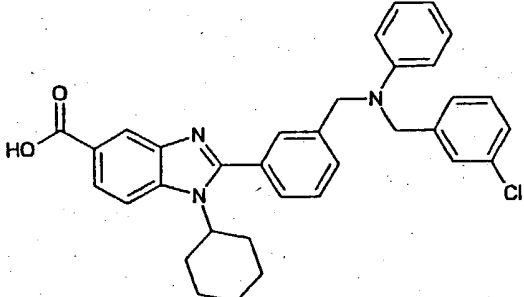
Ex. No.	Formula	MS
1495		580 (M+H)
1496		550 (M+H)
1497		606 (M+H)
1498		580 (M+H)
1499		550 (M+H)

Table 163

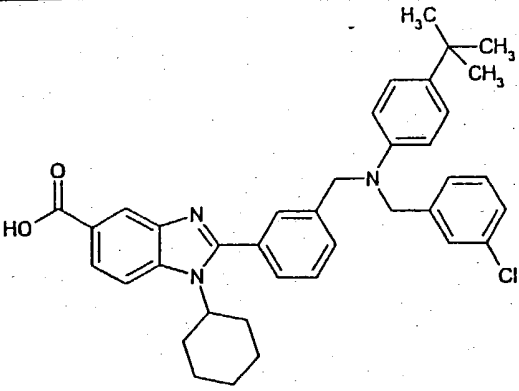
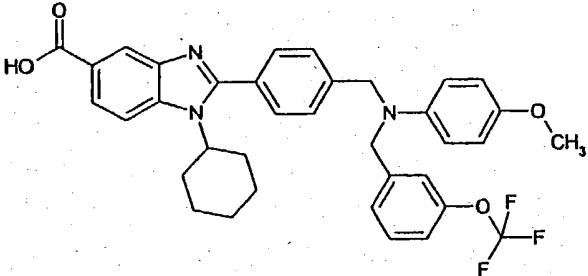
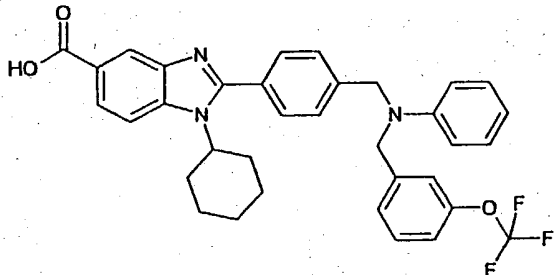
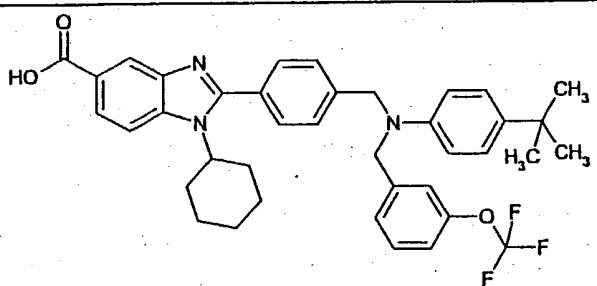
Ex. No.	Formula	MS
1500		606 (M+H)
1501		630 (M+H)
1502		600 (M+H)
1503		656 (M+H)

Table 164

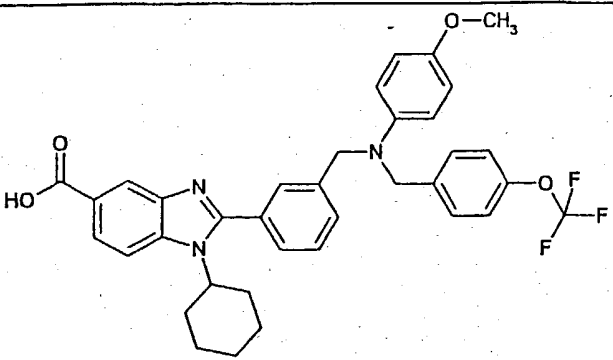
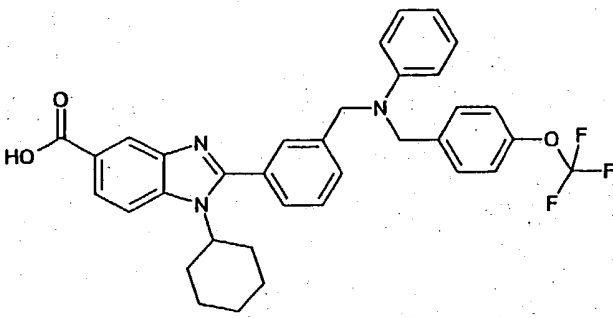
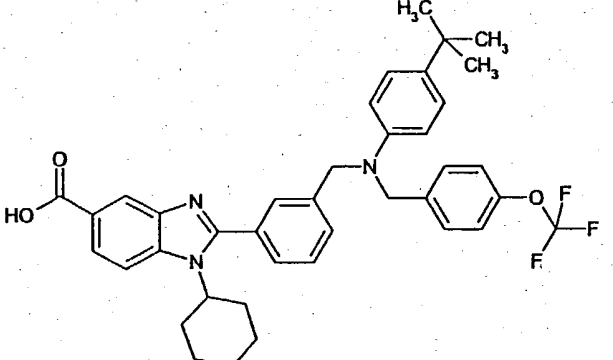
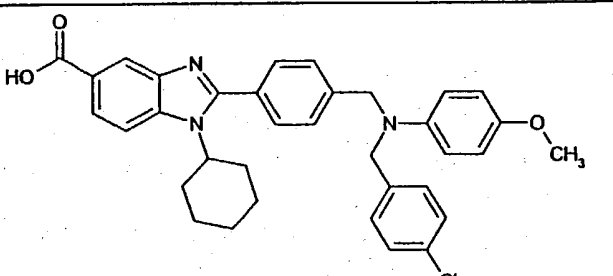
Ex. No.	Formula	MS
1504		630 (M+H)
1505		600 (M+H)
1506		656 (M+H)
1507		580 (M+H)



Table 165

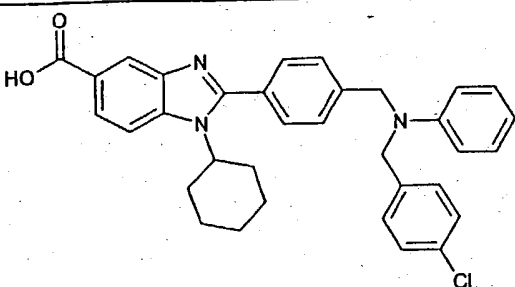
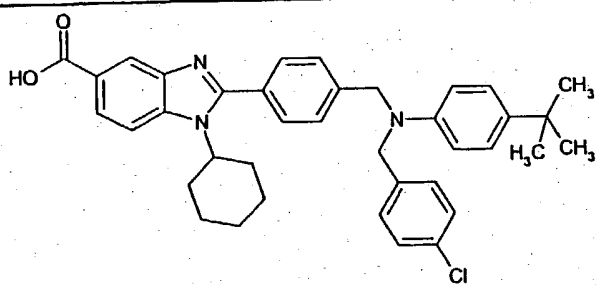
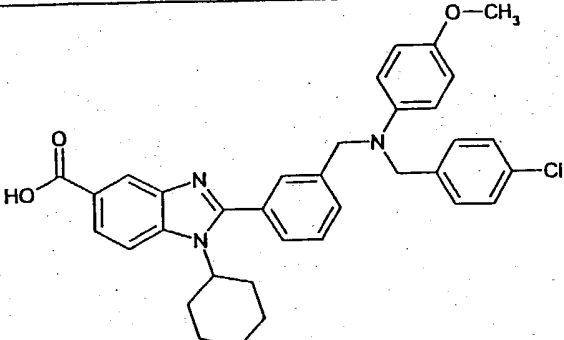
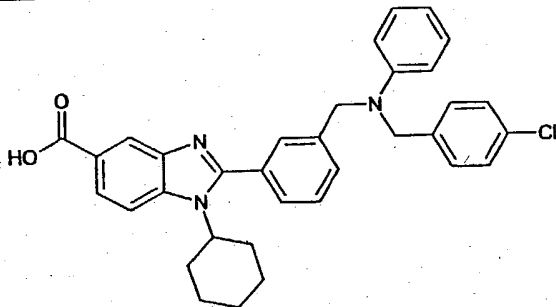
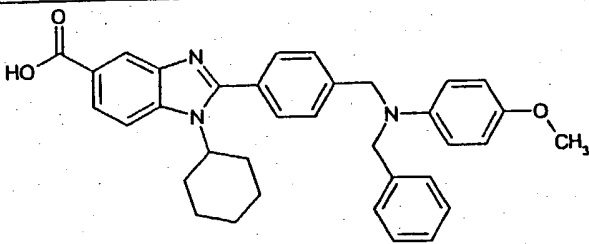
Ex. No.	Formula	MS
1508		550 (M+H)
1509		606 (M+H)
1510		580 (M+H)
1511		550 (M+H)
1512		546 (M+H)

Table 166

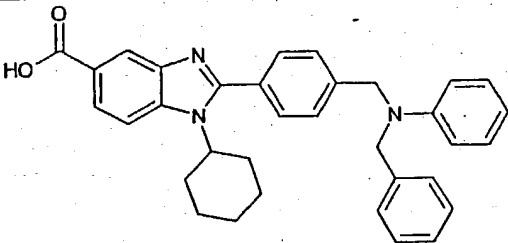
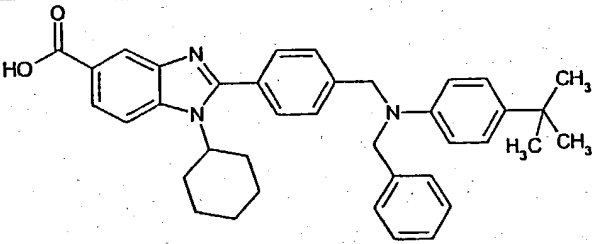
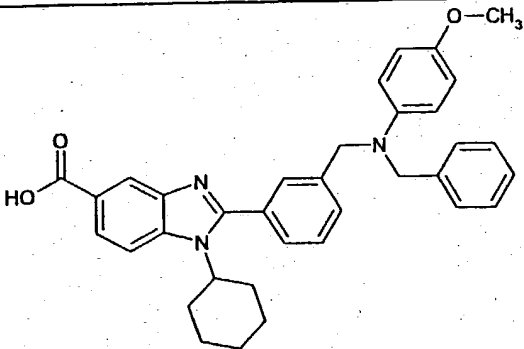
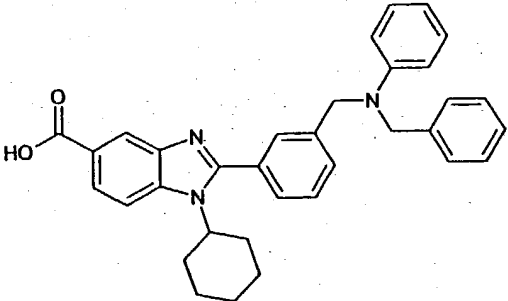
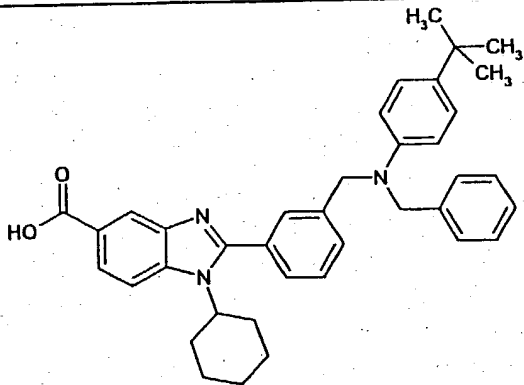
Ex. No.	Formula	MS
1513		516 (M+H)
1514		572 (M+H)
1515		546 (M+H)
1516		516 (M+H)
1517		572 (M+H)

Table 167

Ex. No.	Formula	MS
1518	 <chem>COc1ccc(cc1)N(Cc2ccc(cc2)C(C)(C)C)Cc3ccc(cc3)c4nc5ccccc5n4C(=O)O</chem>	602 (M+H)
1519	 <chem>c1ccccc1N(Cc2ccc(cc2)C(C)(C)C)Cc3ccc(cc3)c4nc5ccccc5n4C(=O)O</chem>	572 (M+H)
1520	 <chem>CC(C)(C)c1ccc(cc1)N(Cc2ccc(cc2)C(C)(C)C)Cc3ccc(cc3)c4nc5ccccc5n4C(=O)O</chem>	628 (M+H)
1521	 <chem>Clc1ccc(cc1)N(Cc2ccc(cc2)C(C)(C)C)Cc3ccc(cc3)c4nc5ccccc5n4C(=O)O</chem>	606 (M+H)

Table 168

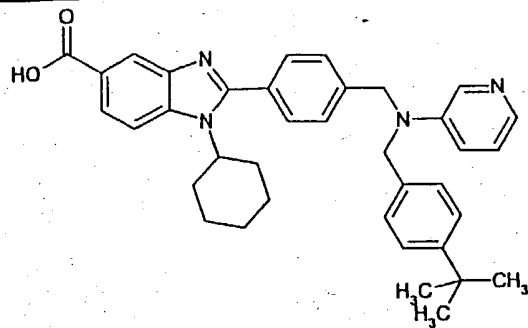
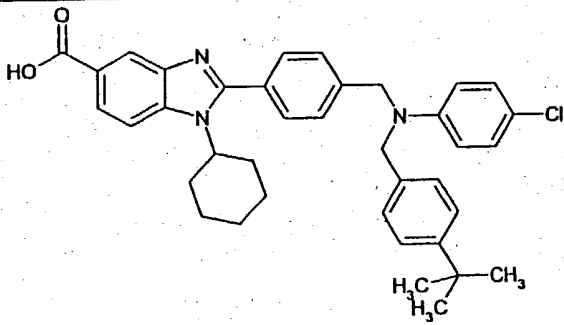
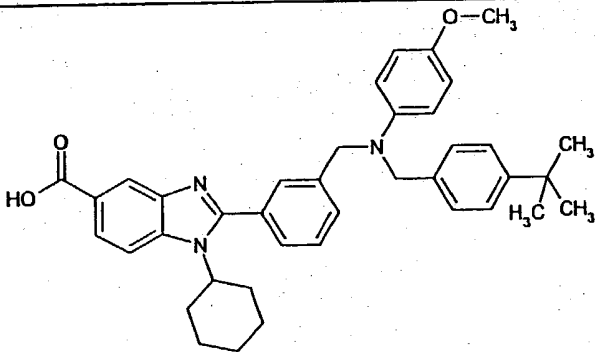
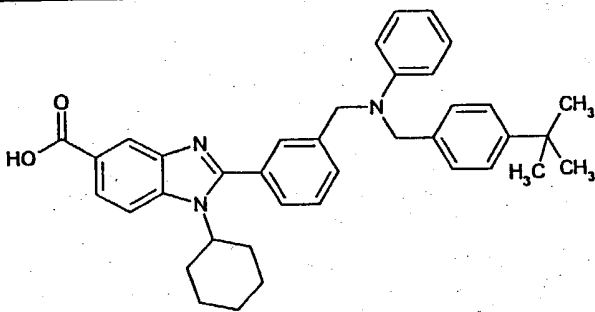
Ex. No.	Formula	MS
1522		573 (M+H)
1523		606 (M+H)
1524		602 (M+H)
1525		572 (M+H)

Table 169

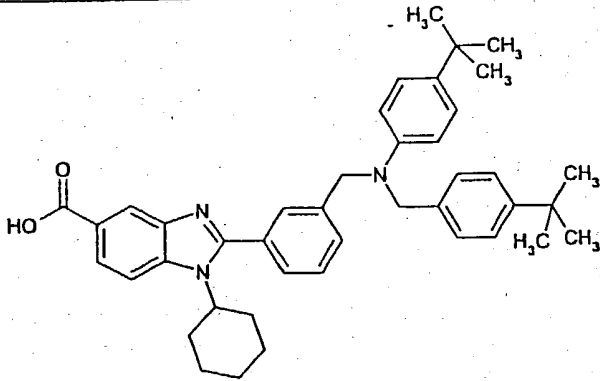
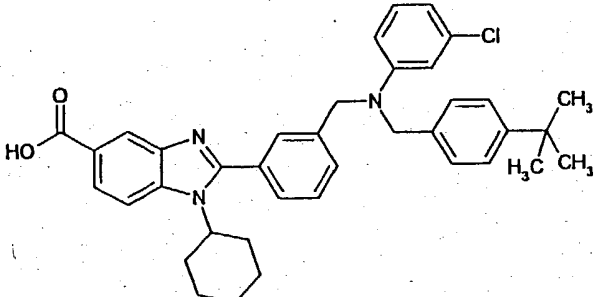
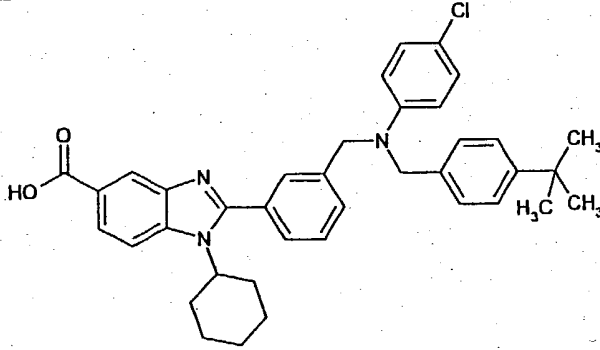
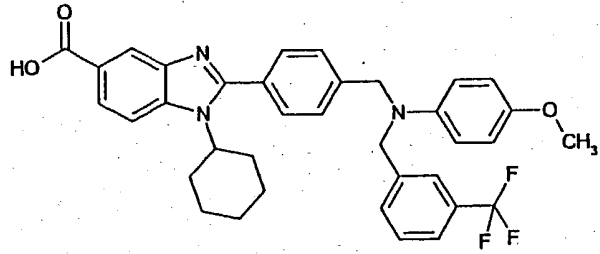
Ex. No.	Formula	MS
1526		628 (M+H)
1527		606 (M+H)
1528		606 (M+H)
1529		614 (M+H)

Table 170

Ex. No.	Formula	MS
1530		584 (M+H)
1531		640 (M+H)
1532		618 (M+H)
1533		614 (M+H)
1534		584 (M+H)

Table 171

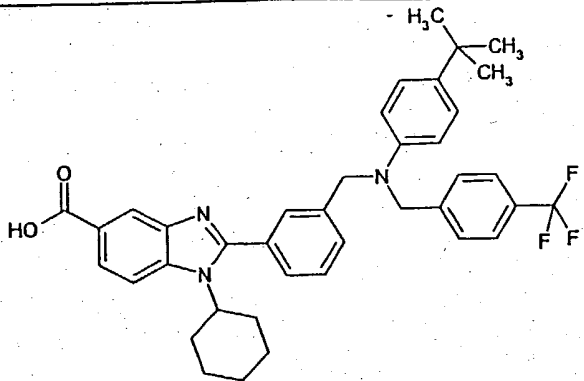
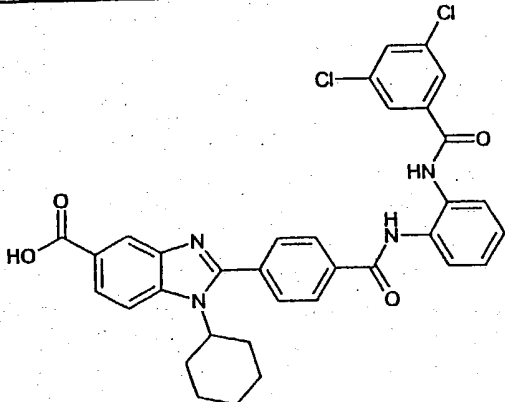
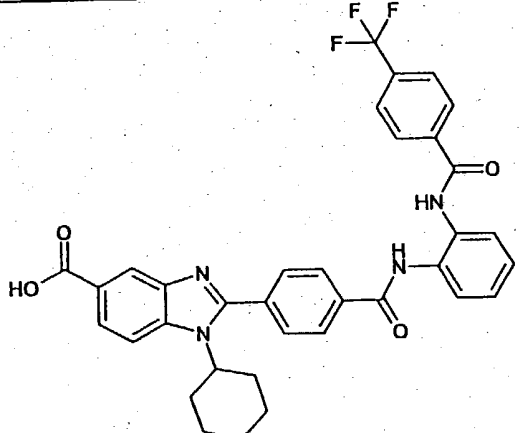
Ex. No.	Formula	MS
1535		640 (M+H)
1536		627 (M+H)
1537		627 (M+H)

Table 172

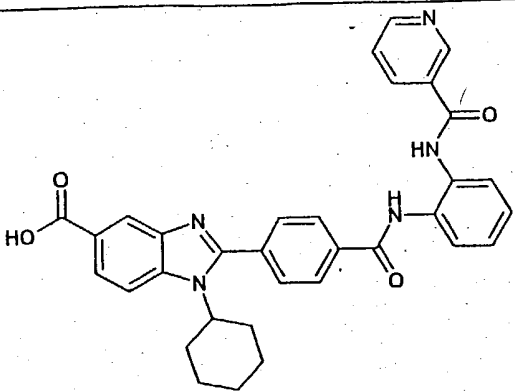
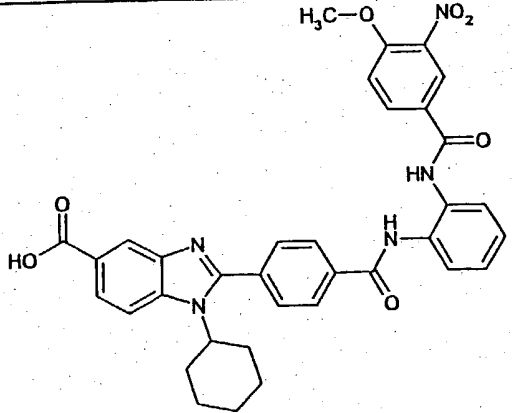
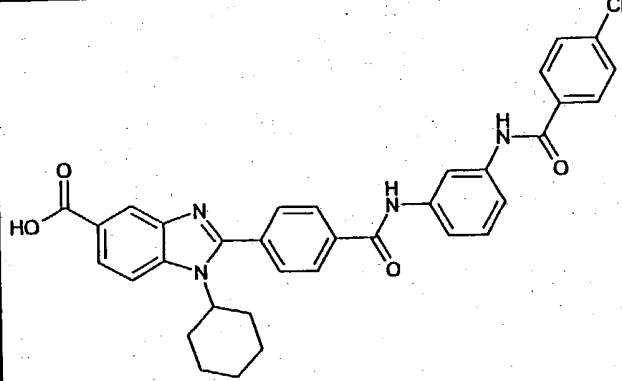
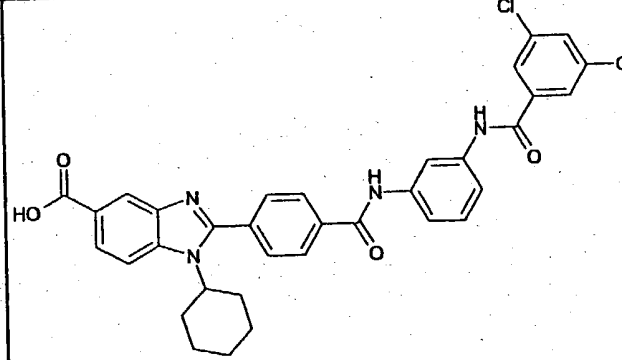
Ex. No.	Formula	MS
1538		560 (M+H)
1539		634 (M+H)
1540		593 (M+H)
1541		627 (M+H)



Table 173

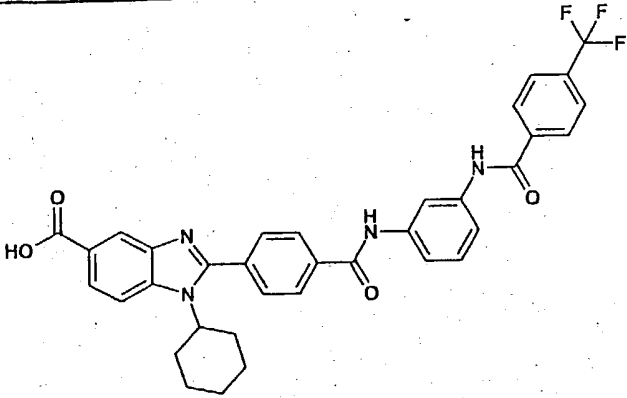
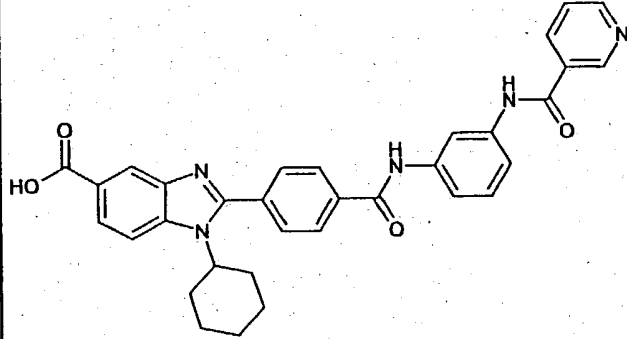
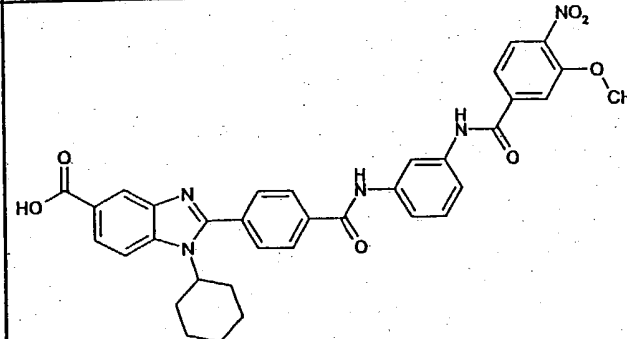
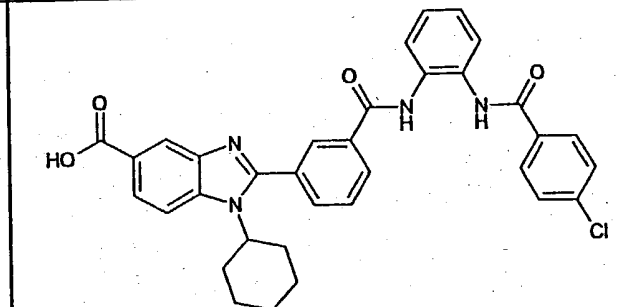
Ex. No.	Formula	MS
1542		627 (M+H)
1543		560 (M+H)
1544		634 (M+H)
1545		593 (M+H)

Table 174

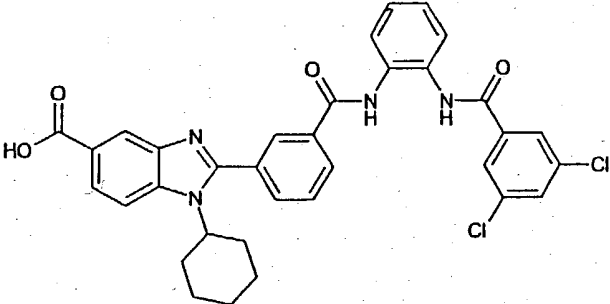
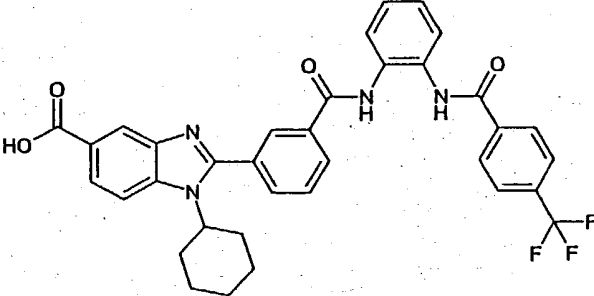
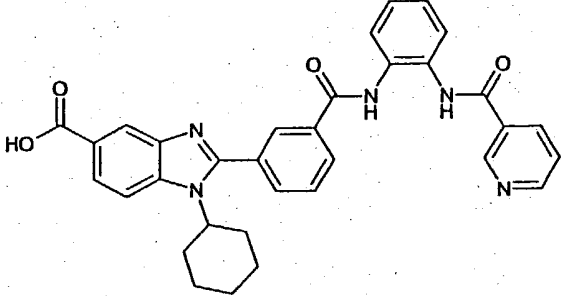
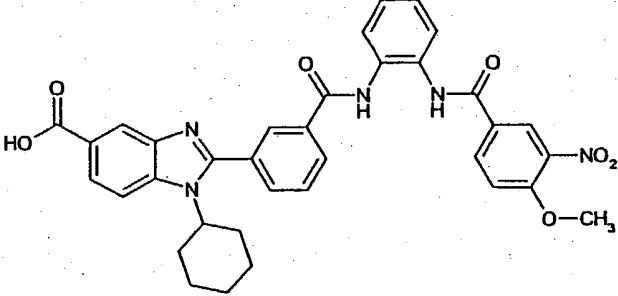
Ex. No.	Formula	MS
1546		627 (M+H)
1547		627 (M+H)
1548		560 (M+H)
1549		634 (M+H)

Table 175

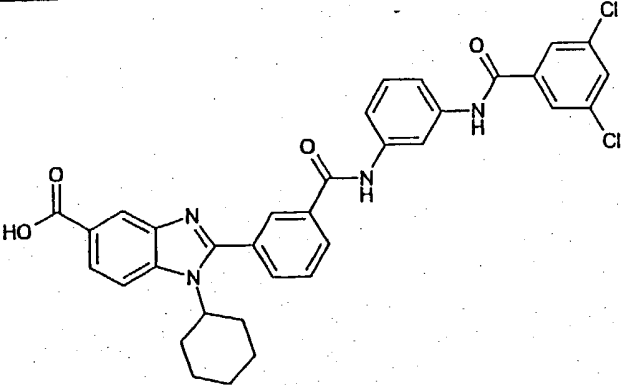
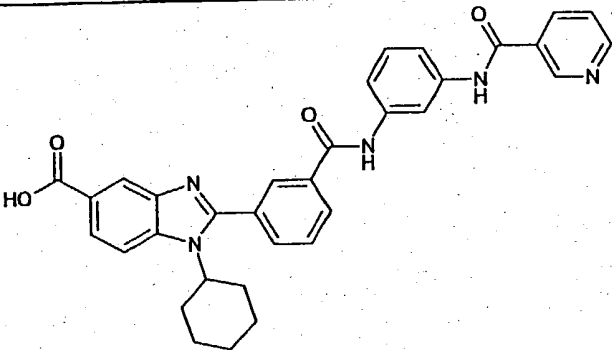
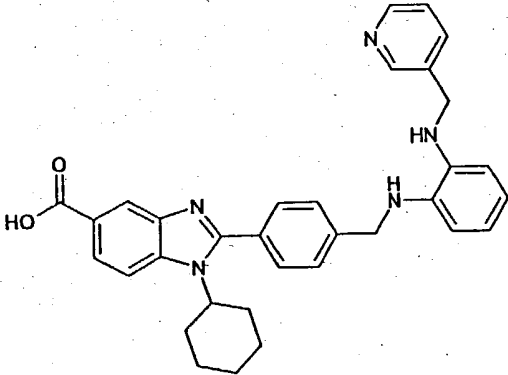
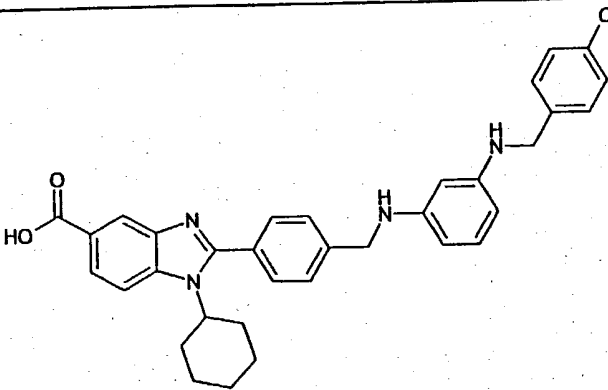
Ex. No.	Formula	MS
1550		627 (M+H)
1551		560 (M+H)
1552		532 (M+H)
1553		565 (M+H)

Table 176

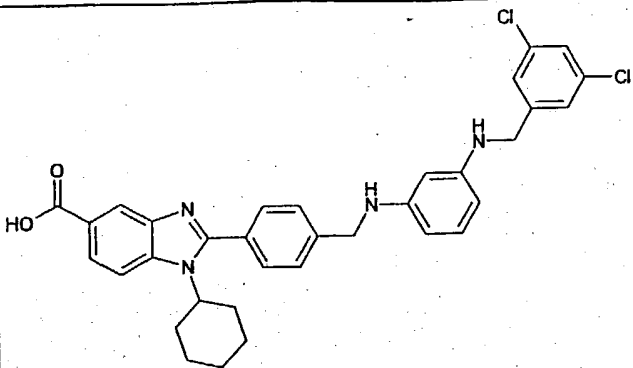
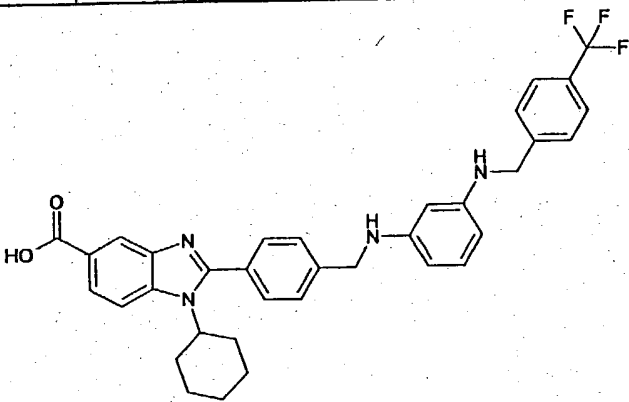
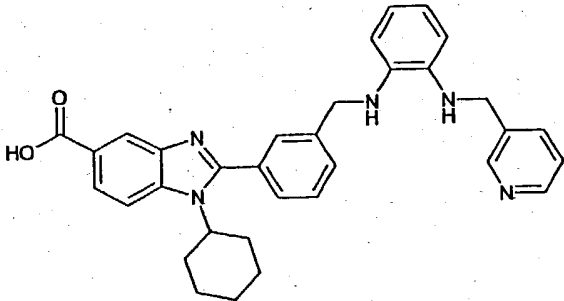
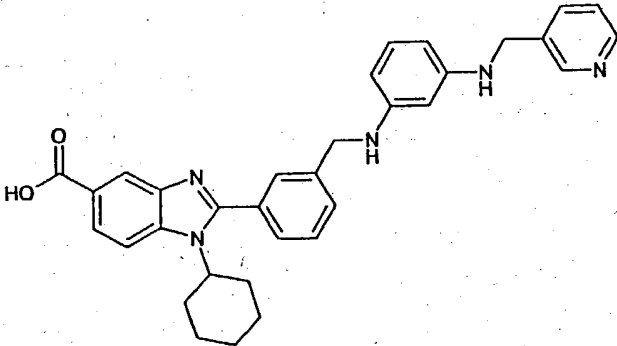
Ex. No.	Formula	MS
1554		599 (M+H)
1555		599 (M+H)
1556		532 (M+H)
1557		532 (M+H)

Table 177

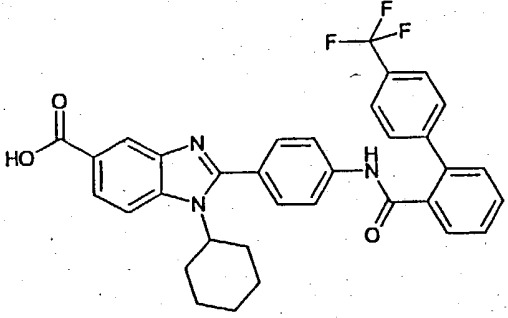
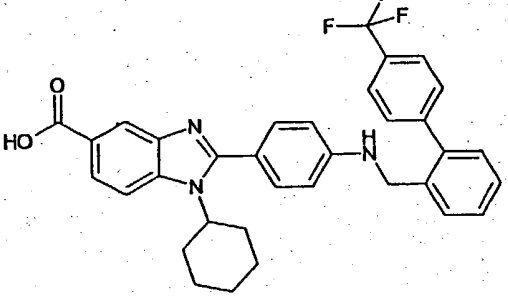
Ex. No.	Formula	MS
1558	 <chem>O=C(O)c1ccc2nc(c1)n(c2)C3CCCCC3c4ccc(cc4)Nc5ccc(cc5)C(F)(F)F</chem>	584 (M+H)
1559	 <chem>O=C(O)c1ccc2nc(c1)n(c2)C3CCCCC3c4ccc(cc4)Nc5ccc(cc5)C(F)(F)F</chem>	570 (M+H)

Table 178

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
2	0.079	67	0.26
6	0.034	68	0.28
9	0.019	70	0.19
11	0.53	71	0.62
12	0.60	77	0.51
17	0.047	81	0.18
20	0.042	82	0.097
26	0.033	83	0.52
30	0.052	85	0.17
43	0.58	86	0.13
44	0.95	87	0.80
45	0.40	88	0.092
46	0.47	89	0.34
47	0.54	90	0.20
48	0.44	91	0.53
49	0.94	93	0.16
50	0.54	94	0.084
51	1.0	96	0.25
54	0.56	97	0.16
55	0.36	98	0.30

Table 179

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
99	0.53	120	0.16
100	0.78	121	0.19
101	0.14	122	0.51
103	0.17	123	0.10
104	0.073	124	0.091
105	0.076	125	0.12
106	0.40	128	0.14
107	0.11	129	0.12
108	0.21	130	0.16
109	0.11	131	0.046
110	0.24	132	0.055
111	0.14	133	0.12
112	0.11	134	0.071
113	0.071	139	0.26
114	0.56	140	0.11
115	0.17	141	0.43
116	0.37	142	0.055
117	0.075	143	0.053
118	0.14	144	0.19
119	0.13	145	0.088

Table 180

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
146	0.043	167	0.033
147	0.31	168	0.078
148	0.038	169	0.15
149	0.15	170	0.048
150	0.24	171	0.050
151	0.20	172	0.10
153	0.19	173	0.14
154	0.076	174	0.030
155	0.53	175	0.29
156	0.23	176	0.053
157	0.16	177	0.077
158	0.11	178	0.052
159	0.13	179	0.63
160	0.24	180	0.11
161	0.062	181	0.71
162	0.43	182	0.021
163	0.15	183	0.017
164	0.16	184	0.018
165	0.58	185	0.11
166	0.055	186	0.37



Table 181

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
187	0.056	207	0.081
188	0.038	208	0.039
189	0.017	209	0.12
190	0.020	210	0.31
191	0.43	211	0.059
192	0.22	212	0.23
193	0.13	213	0.10
194	0.52	214	0.059
195	0.023	215	0.078
196	0.20	216	0.084
197	0.11	217	0.058
198	0.044	218	0.033
199	0.11	219	0.13
200	0.10	220	0.073
201	0.14	221	0.058
202	0.095	222	0.041
203	0.063	223	0.21
204	0.16	225	0.014
205	0.077	227	0.045
206	0.05	228	0.18

Table 182

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]
229	0.022	257	0.074
230	0.17	259	0.10
231	0.073	260	0.27
232	0.015	262	0.013
233	0.028	263	0.035
234	0.022	264	<0.01
235	0.036	265	0.014
236	0.075	266	0.018
237	0.015	267	0.014
238	0.19	268	0.012
239	0.17	269	0.013
240	0.055	270	0.012
248	0.012	271	0.024
249	0.022	272	0.066
250	0.018	273	0.041
252	0.32	276	0.023
253	0.65	279	0.017
254	0.038	280	0.016
255	0.038	281	0.052
256	0.079	282	0.019

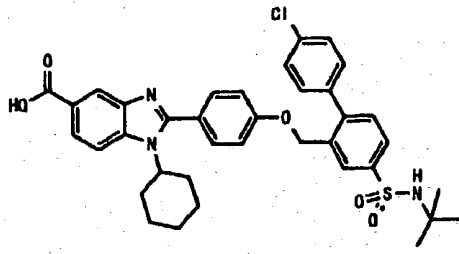
Table 183

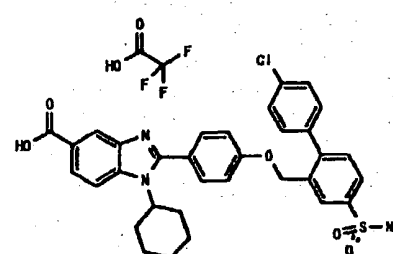
Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]
283	0.014	300	0.045
284	0.014	301	0.017
285	0.012	303	0.10
286	0.014	304	0.017
287	0.012	305	0.01
288	0.013	306	0.013
289	<0.01	307	0.022
290	0.012	308	0.023
291	0.016	311	0.16
292	0.015	312	0.023
293	0.034	313	0.025
294	0.032	314	0.097
295	0.045	315	0.028
296	0.034	316	0.022
297	0.022	317	0.032
298	0.011	318	0.012
299	0.018	319	0.030

Table 184

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]
320	0.036	328	0.015
321	0.015	329	0.047
322	0.016	330	0.011
323	0.018	331	0.017
324	0.027	332	0.023
325	0.019	333	0.016
326	0.018	334	0.016
327	0.019	335	0.013

Table 185

Example No.	249	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.02(1H, d, J=1.5Hz), 8.11(1H, d, J=1.8Hz), 7.96-7.81(3H, m), 7.67(1H, s), 7.61-7.49(6H, m), 7.08(2H, d, J=8.6Hz), 5.19(2H, s), 4.25(1H, m), 2.38-2.17(2H, m), 1.96-1.78(4H, m), 1.70-1.56(1H, m), 1.46-1.16(3H, m), 1.11(9H, s)
Purity	>90% (NMR)	
MS	672(M+1)	

Example No.	250	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.25(1H, d, J=1.5Hz), 8.16-8.08(2H, m), 7.99-7.88(2H, m), 7.66(2H, d, J=8.6Hz), 7.60-7.48(5H, m), 7.19(2H, d, J=8.6Hz), 5.17(2H, s), 4.31(1H, m), 2.39-2.20(2H, m), 2.04-1.79(4H, m), 1.72-1.60(1H, m), 1.50-1.18(3H, m)
Purity	>90% (NMR)	
MS	616(M+1)	

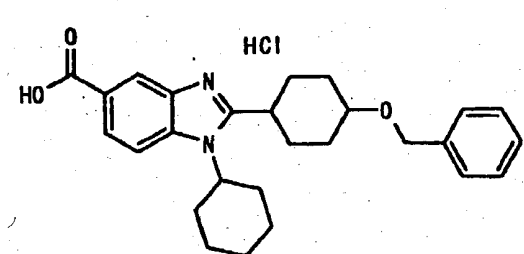
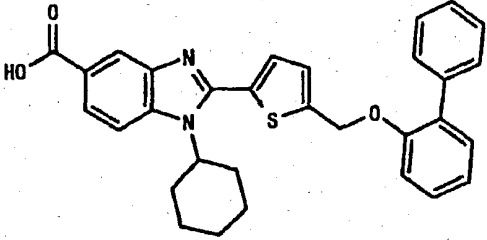
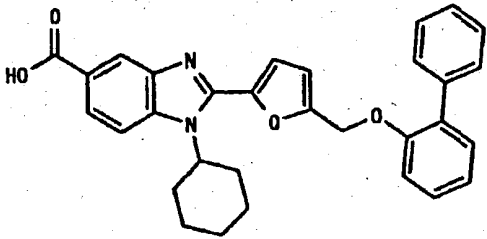
Example No.	251	1H NMR(δ) ppm
		300MHz, DMSO-d6 cis and trans mixture 8.13 and 8.11 (total 1H, each s), 7.90-7.74(2H, m), 7.42-7.22(5H, m), 4.56 and 4.52 (total 2H, each s), 4.42(1H, brs), 3.78-3.06(2H, m) 2.33-1.33(18H, m)
Purity	>90% (NMR)	
MS	433(M+1)	

Table 186

Example No.	252	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.20 (1H, d, J=1.5Hz), 7.96 (1H, d, J=8.6Hz), 7.84 (1H, dd, J=8.6, 1.5Hz), 7.54 (2H, d, J=6.9Hz), 7.48-7.26 (8H, m), 7.09 (1H, t, J=7.3Hz), 5.43 (2H, s), 4.06 (1H, m), 2.40-2.20 (2H, m), 2.01-1.80 (4H, m), 1.75-1.64 (1H, m), 1.51-1.28 (3H, m)
Purity	> 90% (NMR)	
MS	509 (M+1)	

Example No.	253	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.21 (1H, d, J=1.5Hz), 7.93 (1H, d, J=8.7Hz), 7.85 (1H, dd, J=8.4, 1.5Hz), 7.54-7.47 (2H, m), 7.40-7.24 (6H, m), 7.15 (1H, d, J=3.6Hz), 7.11-7.05 (1H, m), 6.81 (1H, d, J=3.6Hz), 5.26 (2H, s), 4.96 (1H, m), 2.32-2.13 (2H, m), 1.95-1.72 (4H, m), 1.68-1.55 (1H, m), 1.43-1.18 (3H, m)
Purity	> 90% (NMR)	
MS	493 (M+1)	

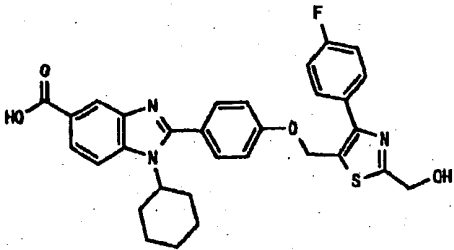
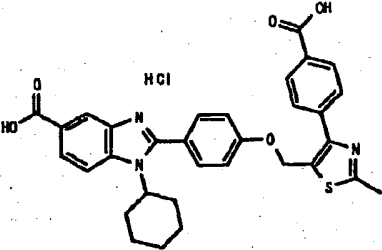
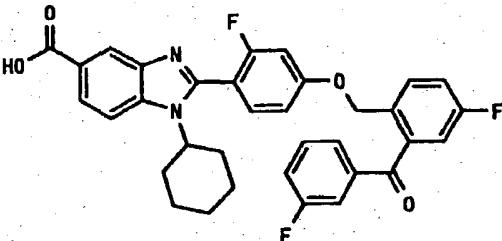
Example No.	254	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.25 (1H, s), 8.02 (1H, d, J=8.7Hz), 7.90 (1H, dd, J=8.4, 1.4Hz), 7.80-7.71 (2H, m), 7.67 (2H, d, J=8.7Hz), 7.33 (2H, t, J=8.7Hz), 7.26 (2H, d, J=8.7Hz), 5.46 (2H, s), 4.78 (2H, s), 4.31 (1H, m), 2.39-2.19 (2H, m), 2.03-1.79 (4H, m), 1.71-1.59 (1H, m), 1.50-1.17 (3H, m)
Purity	> 90% (NMR)	
MS	558 (M+1)	

Table 187

Example No.	255	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.34 (1H, s), 8.32 (1H, d, J=8.8 Hz), 8.09-8.03 (3H, m), 7.83 (2H, d, J=8.3 Hz), 7.79 (2H, d, J=8.8 Hz), 7.36 (2H, d, J=8.8 Hz), 5.54 (2H, s), 4.38 (1H, m), 2.74 (3H, s), 2.40-2.18 (2H, m), 2.13-1.96 (2H, m), 1.93-1.78 (2H, m), 1.73-1.57 (1H, m), 1.55-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	568 (M+1)	

Example No.	256	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.67 (1H, brs), 8.23 (1H, s), 7.94 and 7.87 (2H, ABq, J=8.6 Hz), 7.79 (1H, dd, J=8.7, 5.4 Hz), 7.62-7.41 (7H, m), 6.80 (1H, dd, J=11.9, 2.3 Hz), 6.69 (1H, dd, J=8.1, 2.1 Hz), 5.20 (2H, s), 3.93 (1H, brt, J=15.3 Hz), 2.30-2.11 (2H, brm), 1.88-1.74 (4H, brm), 1.64-1.58 (1H, brm), 1.41-1.14 (3H, brm)
Purity	> 90% (NMR)	
MS	585 (M+1)	

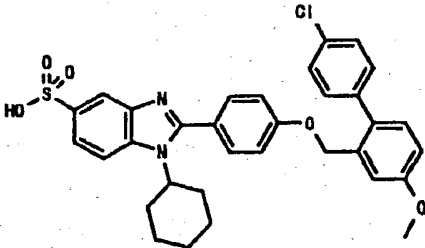
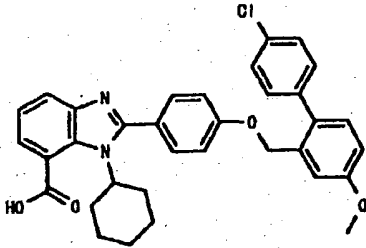
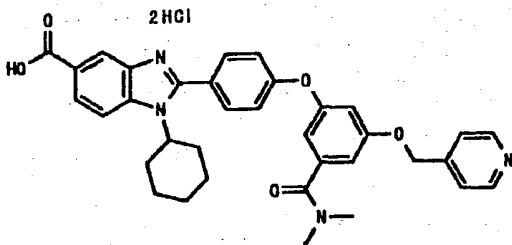
Example No.	257	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.19 (1H, d, J=8.7 Hz), 7.93 (1H, s), 7.83-7.71 (3H, m), 7.50-7.39 (4H, m), 7.34-7.10 (4H, m), 7.06 (1H, dd, J=8.4, 2.9 Hz), 5.09 (2H, s), 4.34 (1H, m), 3.82 (3H, s), 2.39-2.19 (2H, m), 2.11-1.98 (2H, m), 1.94-1.79 (2H, m), 1.74-1.58 (1H, m), 1.52-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	603 (M+1)	

Table 188

Example No.	258	1H NMR(δ) ppm
		300MHz, DMSO-d6 7.79(1H, d, J=6.7Hz), 7.56(1H, d, J=7.5Hz), 7.49(2H, d, J=8.6Hz), 7.42(4H, s), 7.32-7.23(3H, m), 7.09-7.03(3H, m), 5.02(2H, s), 4.46(1H, m), 3.82(3H, s), 1.95-1.83(2H, m), 1.75-1.44(5H, m), 1.30-1.10(2H, m), 0.89-0.71(1H, m)
Purity	> 90% (NMR)	
MS	567 (M+1)	

Example No.	259	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.93(2H, d, J=6.6Hz), 8.36(1H, s), 8.28(1H, d, J=8.7Hz), 8.10-8.03(3H, m), 7.85(2H, d, J=8.7Hz), 7.33(2H, d, J=8.7Hz), 7.23(1H, s), 7.23(1H, s), 6.81(1H, s), 5.56(2H, s), 4.39(1H, m), 2.97, 2.92(6H, s), 2.40-2.18(2H, m), 2.16-1.95(2H, m), 1.90-1.75(2H, m), 1.70-1.55(1H, m), 1.50-1.15(3H, m)
Purity	> 90% (NMR)	
MS	591 (M+1)	

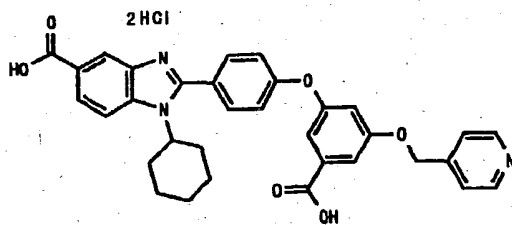
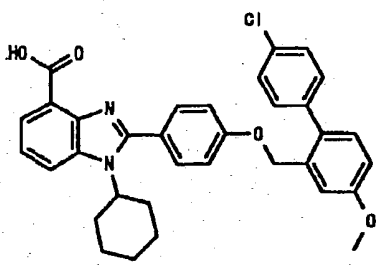
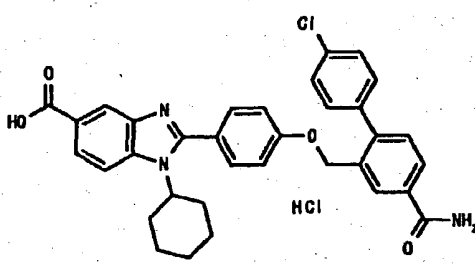
Example No.	260	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.93(2H, d, J=6.3Hz), 8.35(1H, s), 8.26(1H, d, J=8.7Hz), 8.09-8.02(3H, m), 7.86(2H, d, J=8.7Hz), 7.50(1H, s), 7.35(2H, d, J=8.4Hz), 7.24(2H, d, J=7.8Hz), 5.60(2H, s), 4.39(1H, m), 2.50-2.18(2H, m), 2.15-1.95(2H, m), 1.90-1.75(2H, m), 1.70-1.55(1H, m), 1.50-1.10(3H, m)
Purity	> 90% (NMR)	
MS	564 (M+1)	



Table 189

Example No.	261	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.22 (1H, d, J=7.8Hz), 7.85 (1H, d, J=6.7Hz), 7.63 (2H, d, J=9.0Hz), 7.51-7.38 (5H, m), 7.29 (1H, d, J=8.3Hz), 7.23 (1H, d, J=3.0Hz), 7.06 (2H, d, J=9.0Hz), 7.06 (1H, dd, J=8.6, 3.0Hz), 5.05 (2H, s), 4.41-4.25 (1H, m), 3.83 (3H, s), 2.40-2.20 (2H, m), 2.03-1.78 (4H, m), 1.72-1.57 (1H, m), 1.50-1.18 (3H, m)
Purity	> 90 % (NMR)	
MS	567 (M+1)	

Example No.	262	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.29 (1H, d, J=1.5Hz), 8.26 (1H, d, J=9.0Hz), 8.19 (1H, d, J=1.8Hz), 8.13 (1H, brs), 8.08-7.96 (2H, m), 7.73 (2H, d, J=9.0Hz), 7.57-7.43 (6H, m), 7.24 (2H, d, J=9.0Hz), 5.14 (2H, s), 4.36 (1H, m), 2.38-2.18 (2H, m), 2.12-1.97 (2H, m), 1.93-1.80 (2H, m), 1.73-1.58 (1H, m), 1.52-1.20 (3H, m)
Purity	> 90 % (NMR)	
MS	580 (M+1)	

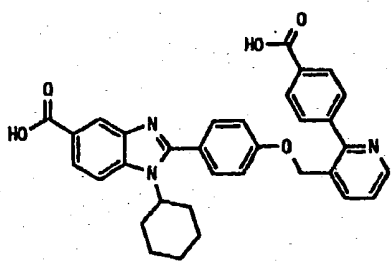
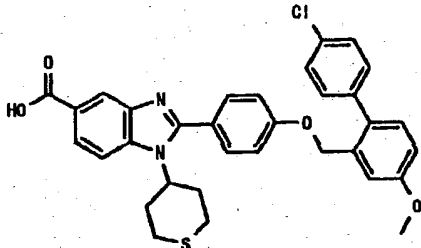
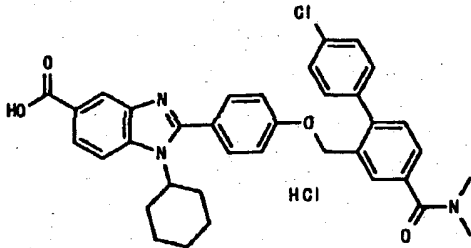
Example No.	263	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.85 (1H, brs), 8.72 (1H, d, J=4.8Hz), 8.22 (1H, s), 8.14 (1H, d, J=6.3Hz), 8.03 and 7.76 (4H, ABq, J=8.6Hz), 7.93a and 7.85 (2H, A'B'q, J=8.6Hz), 7.60 and 7.15 (4H, A''B''q, J=8.7Hz), 7.55 (1H, dd, J=6.3, 4.8Hz), 5.19 (2H, s), 4.26 (1H, brt, J=12.6Hz), 2.35-2.18 (2H, brm), 1.95-1.77 (4H, brm), 1.70-1.60 (1H, brm), 1.45-1.15 (3H, brm)
Purity	> 90 % (NMR)	
MS	548 (M+1)	

Table 190

Example No.	264	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, d, J=1.0Hz), 7.92 (1H, dd, J=8.7, 1.0Hz), 7.87 (1H, d, J=8.7Hz), 7.60 (2H, d, J=8.6Hz), 7.47 (2H, d, J=8.7Hz), 7.44 (2H, d, J=8.7Hz), 7.30 (1H, d, J=8.3Hz), 7.23 (1H, d, J=2.6Hz), 7.11 (2H, d, J=8.7Hz), 7.06 (1H, dd, J=8.7, 2.6Hz), 5.04 (2H, s), 4.36 (1H, m), 3.83 (3H, s), 2.80-2.70 (4H, m), 2.60-2.40 (2H, m), 2.30-2.20 (2H, m)
Purity	> 90% (NMR)	
MS	586, 588 (M+1)	

Example No.	265	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, d, J=1.5Hz), 8.25 (1H, d, J=9.1Hz), 8.03 (1H, dd, J=8.7, 1.5Hz), 7.76-7.96 (3H, m), 7.55-7.49 (5H, m), 7.42 (1H, d, J=7.6Hz), 7.23 (2H, d, J=8.7Hz), 5.15 (2H, s), 4.35 (1H, m), 3.01 (3H, s), 2.97 (3H, s), 2.37-2.20 (2H, m), 2.09-1.97 (2H, m), 1.94-1.81 (2H, m), 1.72-1.60 (1H, m), 1.50-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	608 (M+1)	

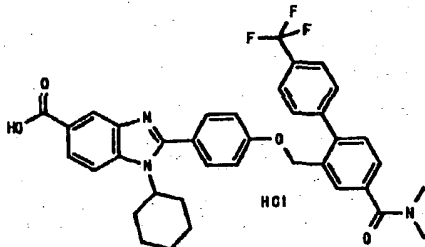
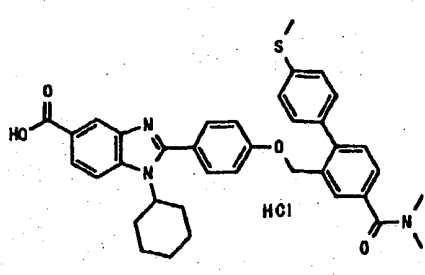
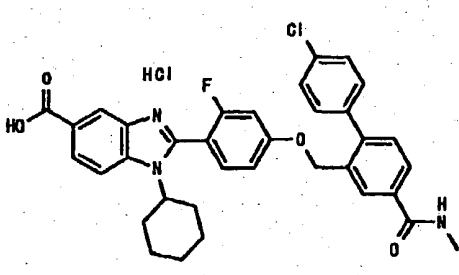
Example No.	266	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, d, J=1.5Hz), 8.20 (1H, d, J=9.0Hz), 8.00 (1H, dd, J=8.6, 1.5Hz), 7.82 (2H, d, J=8.2Hz), 7.76-7.65 (5H, m), 7.56 (1H, dd, J=7.9, 1.8Hz), 7.47 (1H, d, J=7.5Hz), 7.20 (2H, d, J=8.6Hz), 5.16 (2H, s), 4.32 (1H, m), 3.02 (3H, s), 2.98 (3H, s), 2.38-2.19 (2H, m), 2.07-1.95 (2H, m), 1.93-1.80 (2H, m), 1.72-1.58 (1H, m), 1.52-1.18 (3H, m)
Purity	> 90% (NMR)	
MS	642 (M+1)	

Table 191

Example No.	267	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.34 (2H, m), 8.03 (1H, d, J=8.3Hz), 7.77-7.68 (3H, m), 7.54-7.40 (4H, m), 7.33 (2H, d, J=8.6Hz), 7.24 (2H, d, J=9.0Hz), 5.16 (2H, s), 4.36 (1H, m), 3.01 (3H, s), 2.97 (3H, s), 2.40-2.20 (2H, m), 2.11-1.97 (2H, m), 1.93-1.81 (2H, m), 1.71-1.60 (1H, m), 1.50-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	620 (M+1)	

Example No.	268	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.67-8.59 (1H, m), 8.30 (1H, s), 8.13-8.20 (2H, m), 8.02-7.92 (2H, m), 7.65 (1H, t, J=8.3Hz), 7.56-7.45 (5H, m), 7.18 (1H, dd, J=12.0, 2.2Hz), 7.05 (1H, dd, J=8.6, 2.2Hz), 5.14 (2H, s), 4.09 (1H, m), 2.82 (3H, d, J=4.5Hz), 2.34-2.12 (2H, m), 1.99-1.79 (4H, m), 1.71-1.59 (1H, m), 1.49-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	612 (M+1)	

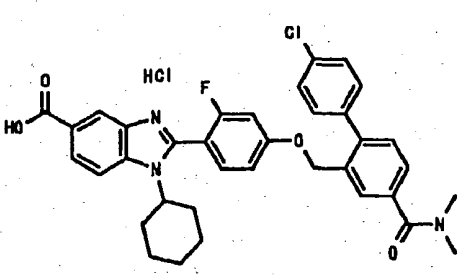
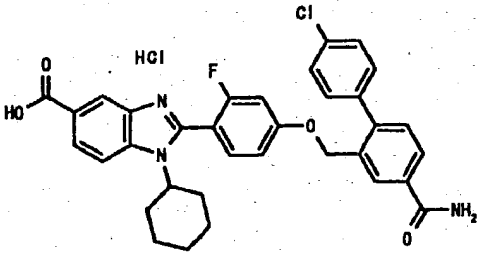
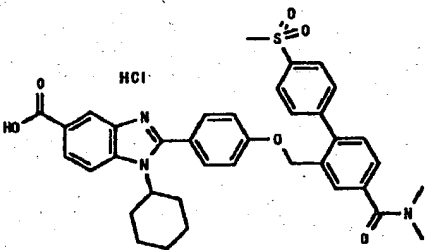
Example No.	269	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.13 (1H, d, J=9.0Hz), 7.97 (1H, dd, J=8.6, 1.5Hz), 7.71 (1H, d, J=1.8Hz), 7.63 (1H, t, J=8.2Hz), 7.56-7.41 (6H, m), 7.17 (1H, dd, J=12.0, 2.2Hz), 7.03 (1H, dd, J=8.2, 1.8Hz), 5.14 (2H, s), 4.15-4.00 (1H, m), 3.01 (3H, s), 2.98 (3H, s), 2.32-2.13 (2H, m), 1.95-1.79 (4H, m), 1.72-1.59 (1H, m), 1.45-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	626 (M+1)	

Table 192

Example No.	270	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.24 (1H, d, J=1.4Hz), 8.19 (1H, d, J=1.8Hz), 8.11 (1H, br s), 8.02-7.85 (3H, m), 7.60-7.44 (7H, m), 7.10 (1H, dd, J=12.0, 2.1Hz), 6.98 (1H, dd, J=8.4, 2.1Hz), 5.11 (2H, s), 3.98 (1H, m), 2.30-2.12 (2H, m), 1.91-1.73 (4H, m), 1.71-1.58 (1H, m), 1.45-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	598 (M+1)	

Example No.	271	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, d, J=1.5Hz), 8.24 (1H, d, J=8.7Hz), 8.07-7.98 (3H, m), 7.80-7.68 (5H, m), 7.56 (1H, dd, J=8.0, 1.8Hz), 7.47 (1H, d, J=8.0Hz), 7.21 (2H, d, J=8.4Hz), 5.18 (2H, s), 4.34 (1H, m), 3.27 (3H, s), 3.02 (3H, s), 2.98 (3H, s), 2.38-2.18 (2H, m), 2.10-1.95 (2H, m), 1.93-1.79 (2H, m), 1.72-1.59 (1H, m), 1.50-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	652 (M+1)	

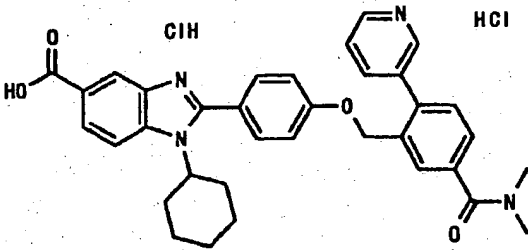
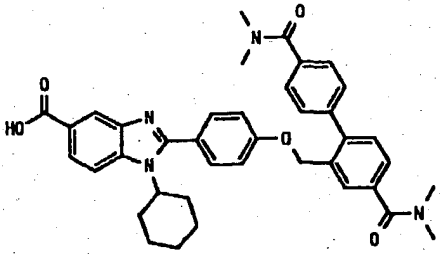
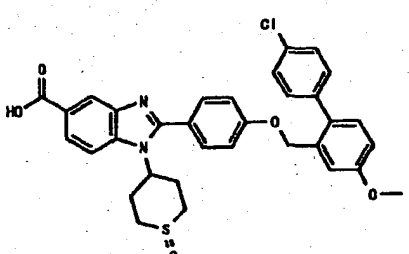
Example No.	272	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.97 (1H, d, J=1.8Hz), 8.85 (1H, d, J=4.7Hz), 8.46 (1H, d, J=8.0Hz), 8.39-8.26 (2H, m), 8.06 (1H, d, J=8.7Hz), 7.99-7.64 (6H, m), 7.24 (2H, d, J=8.7Hz), 5.25 (2H, s), 4.36 (1H, m), 3.03 (3H, s), 2.97 (3H, s), 2.39-2.19 (2H, m), 2.14-1.96 (2H, m), 1.94-1.78 (2H, m), 1.73-1.60 (1H, m), 1.21-1.55 (3H, m)
Purity	> 90% (NMR)	
MS	575 (M+1)	

Table 193

Example No.	273	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.27 (1H, d, J=8.7Hz), 8.05 (1H, d, J=8.7Hz), 7.77-7.67 (3H, m), 7.58-7.48 (6H, m), 7.22 (2H, d, J=8.4Hz), 5.18 (2H, s), 4.35 (1H, brt, J=9.8Hz), 3.06-2.88 (12H, brm), 2.38-2.20 (2H, brm), 2.08-1.96 (2H, brm), 1.90-1.80 (2H, brm), 1.70-1.60 (1H, brm), 1.49-1.22 (3H, brm)
Purity	> 90% (NMR)	
MS	645 (M+1)	

Example No.	274	1H NMR (δ) ppm
		300MHz, DMSO-d6 mixture of cis and trans 8.35, 8.34 (1H, s), 8.15-8.10 (2H, m), 7.79-7.70 (3H, m), 7.49 (2H, d, J=8.7Hz), 7.44 (2H, d, J=8.7Hz), 7.31 (1H, d, J=8.4Hz), 7.25-7.19 (2H, m), 7.07 (1H, d, J=8.5Hz), 5.08 (2H, s), 4.75 (1H, m), 3.83 (3H, s), 3.70-1.90 (8H, m)
Purity	about 80% (NMR)	
MS	601 (M+1)	

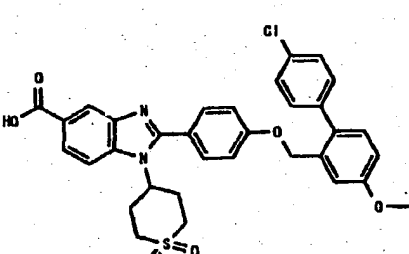
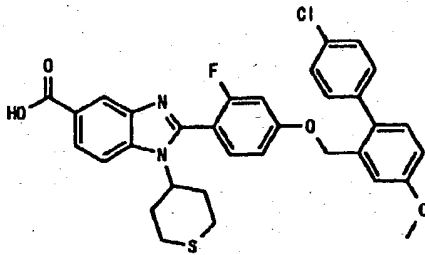
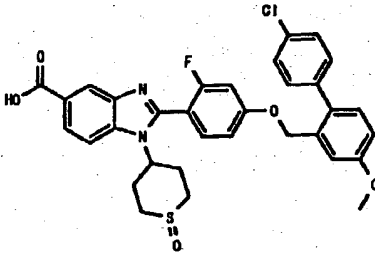
Example No.	275	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.13 (1H, d, J=7.5Hz), 7.93 (1H, d, J=8.8Hz), 7.74 (2H, d, J=8.7Hz), 7.49 (2H, d, J=8.6Hz), 7.44 (2H, d, J=8.6Hz), 7.31 (1H, d, J=8.5Hz), 7.25-7.15 (3H, m), 7.07 (1H, d, J=8.5Hz), 5.08 (2H, s), 4.98 (1H, m), 3.83 (3H, s), 3.65-3.45 (2H, m), 3.30-3.10 (2H, m), 3.00-2.75 (2H, m), 2.60-2.30 (2H, m)
Purity	> 90% (NMR)	
MS	617 (M+1)	

Table 194

Example No.	276	1H NMR(δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.25(1H, s), 7.93and7.87(2H, ABq, J=9.1Hz), 7.55(1H, t, J=8.6Hz), 7.48and7.42(4H, A'B'q, J=8.6Hz), 7.31(1H, d, J=8.5Hz), 7.24(1H, d, J=2.6Hz), 7.09-6.95(3H, m), 5.05(2H, s), 4.11(1H, brt, J=14.0Hz), 3.84(3H, s), 2.83-2.67(4H, brm), 2.50-2.32(2H, brm), 2.21-2.10(2H, brm)
Purity	> 90% (NMR)	
MS	603(M+1)	

Example No.	277	1H NMR(δ) ppm
		300MHz, DMSO-d <sub>6</sub> cis and trans mixture 8.28and8.24(total 1H, each s), 7.94-7.87(1H, m), 7.60-7.41(5H, m), 7.31(1H, d, J=8.5Hz), 7.23-7.21(1H, m), 7.12-7.05(2H, m), 7.00-6.95(1H, m), 5.06and5.05(total 2H, each s), 4.47and4.34(total 1H, each brs), 3.83(3H, s), 3.12-1.76(8H, m)
Purity	> 90% (NMR)	
MS	619(M+1)	

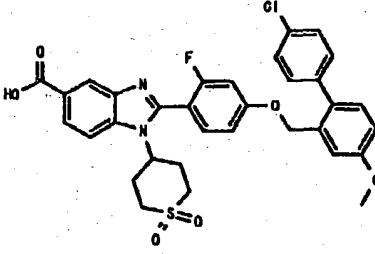
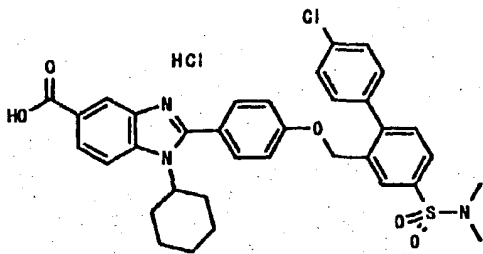
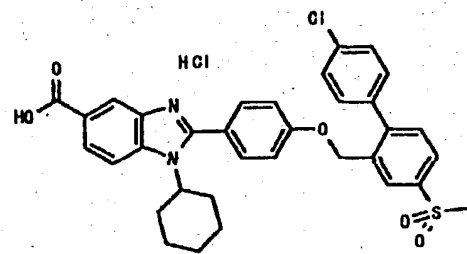
Example No.	278	1H NMR(δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.9(1H, brs), 8.27(1H, s), 7.97and7.74(2H, ABq, J=8.6Hz), 7.58(1H, t, J=8.6Hz), 7.49and7.43(4H, A'B'q, J=8.5Hz), 7.31(1H, d, J=8.5Hz), 7.22(1H, d, J=2.6Hz), 7.13-6.92(3H, m), 5.05(2H, s), 4.67(1H, brt, J=14.2Hz), 3.57-3.40(2H, brm), 3.20-3.05(2H, brm), 2.91-2.70(2H, brm), 2.28-2.11(2H, brm)
Purity	> 90% (NMR)	
MS	635(M+1)	

Table 195

Example No.	279	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 8.23 (1H, d, J=8.7Hz), 8.06-8.00 (2H, m), 7.83 (1H, dd, J=8.0, 1.8Hz), 7.71 (2H, d, J=8.4Hz), 7.64 (1H, d, J=8.0Hz), 7.59-7.54 (4H, m), 7.22 (2H, d, J=8.4Hz), 5.25 (2H, s), 4.33 (1H, m), 2.66 (3H, s), 2.66 (3H, s), 2.37-2.19 (2H, m), 1.93-1.80 (2H, m), 1.70-1.59 (1H, m), 1.47-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	644 (M+1)	

Example No.	280	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.32-8.23 (3H, m), 8.08-8.01 (2H, m), 7.73 (2H, d, J=8.6Hz), 7.65 (1H, d, J=8.2Hz), 7.59-7.51 (4H, m), 7.25 (2H, d, J=8.6Hz), 5.21 (2H, s), 4.34 (1H, m), 3.32 (3H, s), 2.37-2.19 (2H, m), 2.10-1.98 (2H, m), 1.93-1.80 (2H, m), 1.71-1.60 (1H, m), 1.51-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	615 (M+1)	

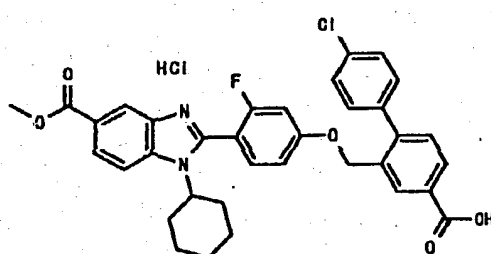
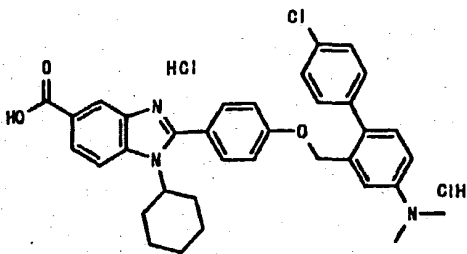
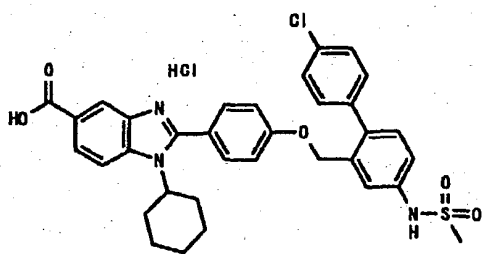
Example No.	281	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, d, J=1.5Hz), 8.24 (1H, s), 8.14 (1H, d, J=8.6Hz), 8.07-7.95 (2H, m), 7.63 (1H, t, J=8.6Hz), 7.57-7.47 (5H, m), 7.16 (1H, dd, J=12.0, 2.2Hz), 7.03 (1H, dd, J=8.6, 2.2Hz), 5.17 (2H, s), 4.06 (1H, m), 3.90 (3H, s), 2.31-2.11 (2H, m), 1.97-1.78 (4H, m), 1.71-1.59 (1H, m), 1.43-1.22 (3H, m)
Purity	> 90% (NMR)	
MS	315	

Table 196

Example No.	282	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.36 (1H, s), 8.35 (1H, d, J=9.3Hz), 8.09 (1H, d, J=9.3Hz), 7.78 (2H, d, J=8.7Hz), 7.48-7.25 (9H, m), 5.09 (2H, s), 4.39 (1H, m), 3.04 (6H, s), 2.40-2.15 (2H, m), 2.10-1.95 (2H, m), 1.90-1.75 (2H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	580 (M+1)	

Example No.	283	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 10.03 (1H, s), 8.33 (1H, s), 8.29 (1H, d, J=8.7Hz), 8.06 (1H, d, J=9.0Hz), 7.74 (2H, d, J=9.0Hz), 7.51-7.42 (5H, m), 7.37-7.30 (2H, m), 7.22 (2H, d, J=8.7Hz), 5.10 (2H, s), 4.37 (1H, m), 3.06 (3H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	630 (M+1)	

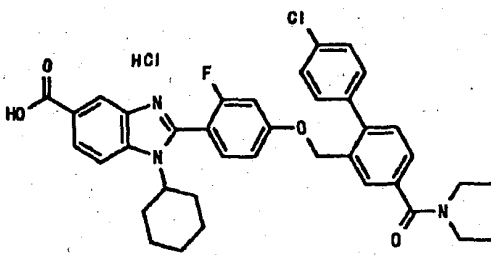
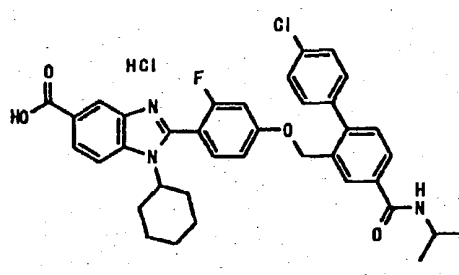
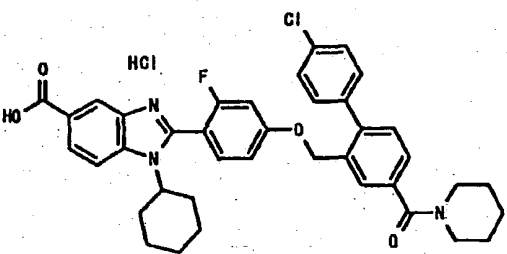
Example No.	284	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 8.14 (1H, d, J=8.7Hz), 7.97 (1H, d, J=8.7Hz), 7.96-7.41 (8H, m), 7.16 (1H, dd, J=12.4, 2.2Hz), 7.03 (1H, dd, J=8.4, 2.2Hz), 5.15 (2H, s), 4.15 (1H, m), 3.54-3.16 (4H, m), 2.33-2.13 (2H, m), 1.97-1.79 (4H, m), 1.70-1.02 (9H, m)
Purity	> 90% (NMR)	
MS	654 (M+1)	



Table 197

Example No.	285	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.37(1H, d, J=7.3Hz), 8.30(1H, s), 8.19-8.12(2H, m), 8.02-7.95(2H, m), 7.65(1H, t, J=8.4Hz), 7.56-7.43(5H, m), 7.18(1H, dd, J=12.0, 1.8Hz), 7.06(1H, dd, J=8.4, 2.1Hz), 5.13(2H, s), 4.22-4.03(2H, m), 2.34-2.13(2H, m), 1.99-1.78(4H, m), 1.72-1.57(1H, m), 1.44-1.14(3H, m), 1.20, 1.18(6H, each s)
Purity	>90% (NMR)	
MS	640 (M+1)	

Example No.	286	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.29(1H, s), 8.13(1H, d, J=8.7Hz), 7.97(1H, dd, J=8.7, 1.4Hz), 7.69-7.40(8H, m), 7.16(1H, dd, J=12.0, 2.2Hz), 7.02(1H, dd, J=8.4, 2.2Hz), 5.15(2H, s), 4.07(1H, m), 3.71-3.23(2H, m), 1.98-1.71(4H, m), 1.71-1.18(10H, m)
Purity	>90% (NMR)	
MS	666 (M+1)	

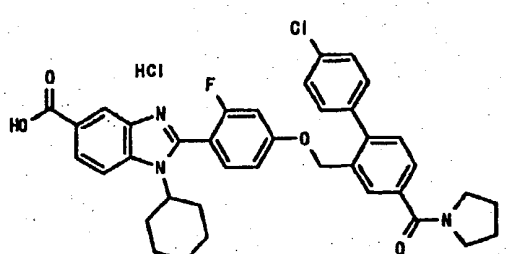
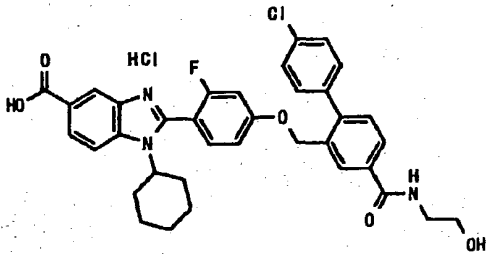
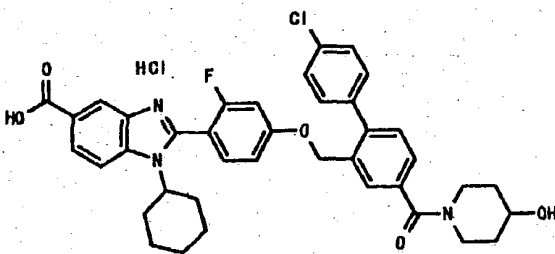
Example No.	287	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.29(1H, s), 8.13(1H, d, J=8.0Hz), 7.97(1H, d, J=8.4Hz), 7.83(1H, s), 7.68-7.41(7H, m), 7.17(1H, d, J=12.0Hz), 7.03(1H, d, J=8.4Hz), 5.15(2H, s), 4.07(1H, m), 3.58-3.41(4H, m), 2.34-2.13(2H, m), 1.97-1.77(8H, m), 1.71-1.58(1H, m), 1.49-1.18(3H, m)
Purity	>90% (NMR)	
MS	652 (M+1)	

Table 198

Example No.	288	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.62 (1H, m), 8.31 (1H, s), 8.22-8.14 (2H, m), 8.99 (2H, d, J=8.7Hz), 7.66 (1H, t, J=7.7Hz), 7.58-7.44 (5H, m), 7.19 (1H, dd, J=8.7, 2.2Hz), 5.14 (2H, s), 4.11 (1H, m), 3.67-3.49 (2H, m), 3.45-3.30 (2H, m), 2.37-2.12 (2H, m), 2.00-1.76 (4H, m), 1.70-1.58 (1H, m), 1.48-1.17 (3H, m)
Purity	> 90% (NMR)	
MS	642 (M+1)	

Example No.	289	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.28 (1H, s), 8.11 (1H, d, J=8.9Hz), 7.96 (1H, d, J=8.9Hz), 7.68 (1H, s), 7.62 (1H, t, J=8.2Hz), 7.55-7.41 (6H, m), 7.15 (1H, d, J=11.7Hz), 7.02 (1H, d, J=8.4Hz), 5.14 (2H, s), 4.12-3.13 (6H, m), 2.30-1.19 (13H, m)
Purity	> 90% (NMR)	
MS	682 (M+1)	

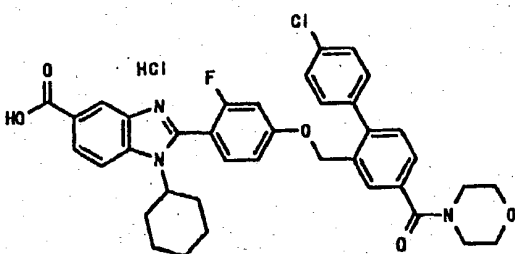
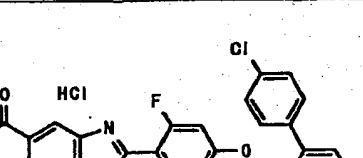
Example No.	290	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.29 (1H, s), 8.15 (1H, d, J=8.6Hz), 7.98 (1H, d, J=8.8Hz), 7.72 (1H, s), 7.64 (1H, t, J=8.8Hz), 7.57-7.43 (6H, m), 7.18 (1H, dd, J=12.1, 2.1Hz), 7.03 (1H, d, J=10.7Hz), 5.12 (2H, s), 4.15-4.01 (1H, m), 3.75-3.33 (8H, m), 2.31-2.14 (2H, m), 1.96-1.78 (4H, m), 1.70-1.58 (1H, m), 1.47-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	668 (M+1)	

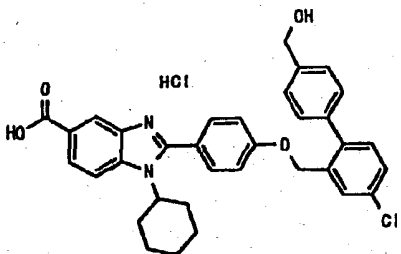
Table 199

Example No.	291	<sup>1</sup> H NMR (δ) ppm
		400MHz, DMSO-d <sub>6</sub> 8.29 (1H, s), 8.14 (1H, d, J=8.9Hz), 7.97 (1H, d, J=8.6Hz), 7.71 (1H, s), 7.63 (1H, t, J=8.2Hz), 7.56-7.42 (6H, m), 7.17 (1H, d, J=12.3Hz), 7.03 (1H, d, J=10.7Hz), 5.14 (2H, s), 4.07 (1H, m), 3.96-3.52 (4H, m), 2.79-2.56 (4H, m), 2.32-2.14 (2H, m), 1.97-1.79 (4H, m), 1.71-1.58 (1H, m), 1.51-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	684 (M+1)	

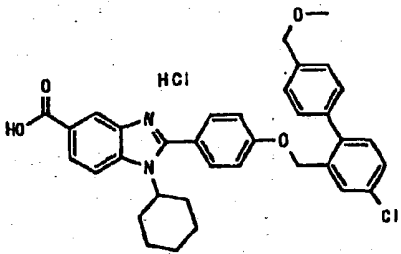
Example No.	292	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9.07-8.99 (1H, m), 8.30 (1H, s), 8.23-8.12 (2H, m), 8.04-7.95 (2H, m), 7.65 (1H, t, J=8.2Hz), 7.60-7.45 (5H, m), 7.19 (1H, dd, J=12.0, 2.6Hz), 7.06 (1H, dd, J=8.6, 2.2Hz), 5.16 (2H, s), 4.18-4.02 (1H, m), 3.97 (2H, d, J=6.0Hz), 2.33-2.14 (2H, m), 1.99-1.79 (4H, m), 1.72-1.59 (1H, m), 1.45-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	656 (M+1)	

Example No.	293	<sup>1</sup> H NMR(δ) ppm
		300MHz, DMSO-d <sub>6</sub> : 8. 21 (1H, s), 7. 94 and 7. 86 (2H, ABq, J=8. 6Hz), 7. 72 (1H, d, J=2. 4Hz), 7. 59 and 7. 11 (4H, A'B'q, J=8. 9Hz), 7. 53 (1H, dd, J=8. 4, 2. 4Hz), 7. 38 (1H, d, J=8. 4Hz), 7. 36 and 7. 32 (4H, A''B''q, J=8. 1Hz), 5. 07 (2H, s), 4. 27 (1H, brt, J=13. 8Hz), 2. 87 (2H, t, J=7. 8Hz), 2. 57 (2H, t, J=7. 8Hz), 2. 35-2. 20 (2H, brm), 1. 96-1. 79 (4H, brm), 1. 68-1. 59 (1H, brm), 1. 47-1. 18 (3H, brm)
Purity	> 90% (NMR)	
MS	637 (M+1)	

Table 200

Example No.	294	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 8.25 and 8.03 (2H, ABq, J=8.9Hz), 7.73 (1H, s), 7.73 (2H, d, J=8.6Hz), 7.55 (1H, dd, J=8.0, 2.3Hz), 7.40 (4H, s), 7.39 (1H, d, J=8.0Hz), 7.23 (2H, d, J=8.6Hz), 5.11 (2H, s), 4.55 (2H, s), 4.36 (1H, brt, J=14.8Hz), 2.37-2.19 (2H, brm), 2.09-1.96 (2H, brm), 1.91-1.79 (2H, brm), 1.71-1.59 (1H, brm), 1.50-1.20 (3H, brm)
Purity	> 90% (NMR)	
MS	567 (M+1)	

Example No.	295	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 8.25 and 8.04 (2H, ABq, J=8.7Hz), 7.74 (1H, s), 7.72 (2H, d, J=8.7Hz), 7.56 (1H, d, J=8.7Hz), 7.48-7.35 (5H, m), 7.22 (2H, d, J=8.7Hz), 5.11 (2H, s), 4.46 (2H, s), 4.35 (1H, brt, J=14.8Hz), 3.31 (3H, s), 2.37-2.17 (2H, brm), 2.07-1.95 (2H, brm), 1.92-1.79 (2H, brm), 1.73-1.56 (1H, brm), 1.52-1.20 (3H, brm)
Purity	> 90% (NMR)	
MS	581 (M+1)	

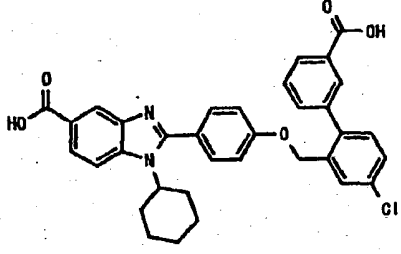
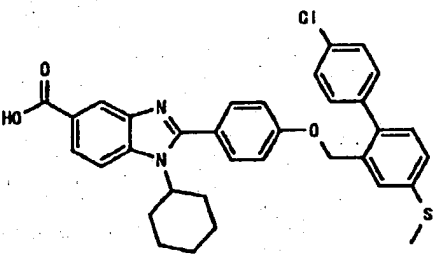
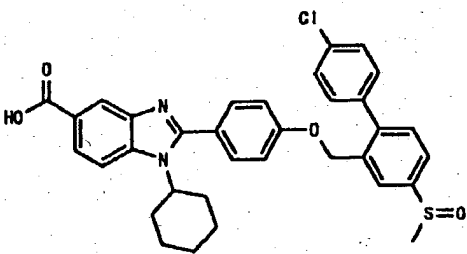
Example No.	296	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.21 (1H, d, J=1.5Hz), 7.98 (1H, d, J=1.2Hz), 7.97-7.91 (2H, m), 7.84 (1H, dd, J=8.7, 1.5Hz), 7.77 (1H, d, J=2.1Hz), 7.70 (1H, d, J=7.5Hz), 7.60-7.54 (4H, m), 7.43 (1H, d, J=8.4Hz), 7.09 (2H, d, J=8.7Hz), 5.05 (2H, s), 4.25 (1H, brt, J=14.8Hz), 2.36-2.18 (2H, brm), 1.95-1.79 (4H, brm), 1.71-1.6 (1H, brm), 1.43-1.18 (3H, brm)
Purity	> 90% (NMR)	
MS	581 (M+1)	

Table 201

Example No.	297	1H NMR (δ) ppm
		300MHz, DMSO-d6 12.7 (1H, brs), 8.21 (1H, s), 7.94 and 7.85 (2H, ABq, J=8.6 Hz), 7.60-7.55 (3H, m), 7.49 and 7.45 (4H, A' B' q, J=8.3Hz , 7.12 (2H, d, J=8.7Hz), 5.0 5 (2H, s), 4.26 (1H, brt, J=13 .0Hz), 2.54 (3H, s), 2.38-2. 20 (2H, brm), 1.97-1.80 (4H, brm), 1.71-1.59 (1H, brm), 1 .47-1.20 (3H, brm)
Purity	> 90% (NMR)	
MS	583 (M+1)	

Example No.	298	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 8.01 (1H, s), 7. 95 and 7.86 (2H, ABq, J=8.6Hz , 7.79 (1H, d, J=7.8Hz), 7.5 8 (3H, t, J=7.5Hz), 7.53 (4H, s), 7.13 (2H, d, 8.7Hz), 5.15 (2H, s), 4.26 (1H, brt, J=13. 8Hz), 2.83 (3H, s), 2.37-2.1 8 (2H, brm), 1.95-1.78 (4H, b rm), 1.70-1.59 (1H, brm), 1. 47-1.17 (3H, brm)
Purity	> 90% (NMR)	
MS	599 (M+1)	

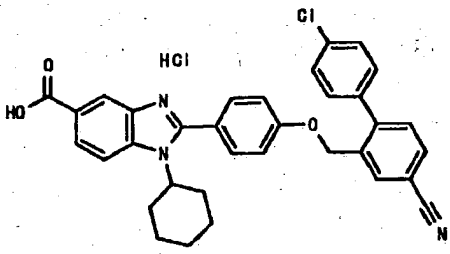
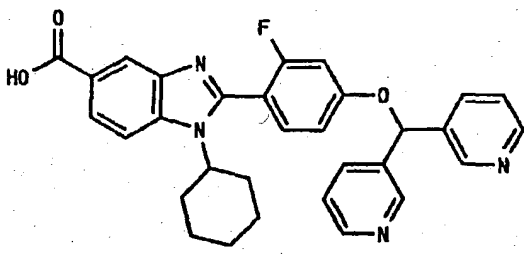
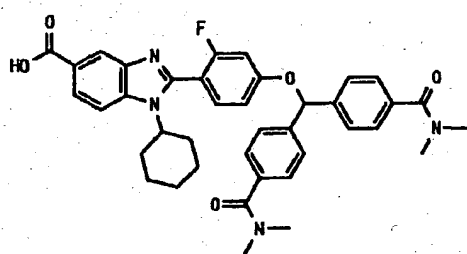
Example No.	299	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.43-8.16 (3H, m), 8.07-7.9 4 (2H, m), 7.72 (2H, d, J=8.6H z), 7.62-7.49 (5H, m), 7.23 (2 H, d, J=8.6Hz), 5.16 (2H, s) , 4.34 (1H, m), 2.39-2.20 (2H , m), 2.10-1.96 (2H, m), 1.93 -1.80 (2H, m), 1.71-1.58 (1H , m), 1.49-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	562 (M+1)	

Table 202

Example No.	300	1H NMR (δ) ppm
		300MHz, DMSO-d6: 2.77 (1H, brs), 8.83 (2H, d, J=1.9Hz), 8.56 (2H, dd, J=4.9, 1.9Hz), 8.22 (1H, d, J=1.5Hz), 7.97 (2H, dt, J=7.9, 1.9Hz), 7.95 (1H, d, J=8.6Hz), 7.87 (1H, dd, J=8.6, 1.5Hz), 7.57 (1H, t, J=8.7Hz), 7.46 (2H, dd, J=7.9, 4.9Hz), 7.26 (1H, dd, J=12.0, 4.9Hz), 7.14 (1H, dd, J=8.8, 2.3Hz), 6.99 (2H, s), 3.94 (1H, brt), 2.26-2.09 (2H, m), 1.87-1.73 (4H, m), 1.67-1.57 (1H, m), 1.42-1.12 (2H, m)
Purity	> 90% (NMR)	
MS	523 (M+1)	

Example No.	301	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.22 (1H, s), 7.95 (1H, d, J=8.7Hz), 7.87 (1H, dd, J=1.5Hz, 9.0Hz), 7.62 (4H, d, J=8.4Hz), 7.55 (1H, t, J=9.0Hz), 7.44 (4H, d, J=8.1Hz), 7.20 (1H, dd, J=2.1Hz, 12.0Hz), 7.11 (1H, dd, J=2.1Hz, 8.7Hz), 6.86 (1H, s), 3.94 (1H, m), 2.96, 2.88 (12H, s), 2.35-2.00 (2H, m), 1.95-1.70 (4H, m), 1.65-1.50 (1H, m), 1.45-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	663 (M+1)	

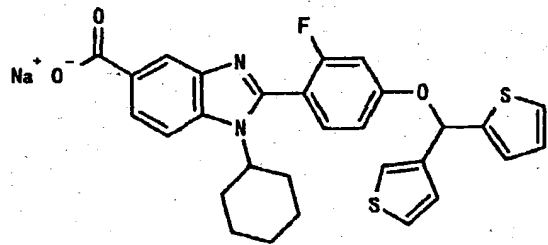
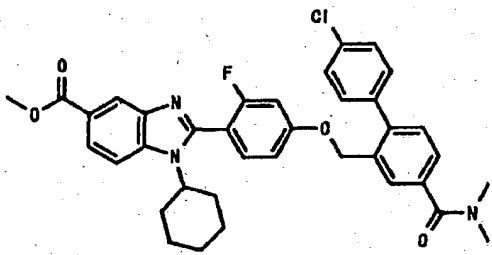
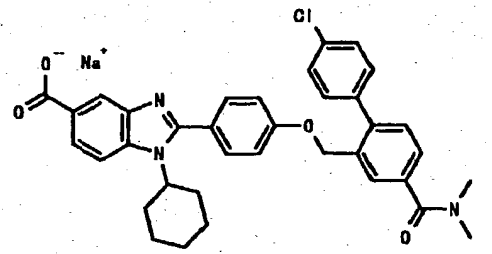
Example No.	302	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.14 (1H, s), 7.88 (1H, d, J=8.4Hz), 7.68 (1H, d, J=8.7Hz), 7.64-7.55 (3H, m), 7.50 (1H, t, J=8.7Hz), 7.22-7.17 (3H, m), 7.11 (1H, s), 7.08-7.00 (2H, m), 3.90 (1H, m), 2.15-2.00 (2H, m), 1.95-1.50 (5H, m), 1.45-1.00 (3H, m)
Purity	> 90% (NMR)	
MS	532 (M+1)	

Table 203

Example No.	303	1H NMR (δ) ppm
		300MHz, CDCl <sub>3</sub> 8.49 (1H, s), 7.98 (1H, dd, J=8.6, 1.5Hz), 7.71 (1H, d, J=1.8Hz), 7.66 (1H, d, J=8.6Hz), 7.55-7.29 (7H, m), 6.80 (1H, dd, J=8.2, 2.2Hz), 6.69 (1H, dd, J=11.2, 2.2Hz), 4.99 (2H, s), 4.10-3.92 (1H, m), 3.95 (3H, s), 3.15 (3H, s), 3.06 (3H, s), 2.31-2.14 (2H, m), 2.04-1.86 (4H, m), 1.81-1.71 (1H, m), 1.41-1.21 (3H, m)
Purity	> 90% (NMR)	
MS	640 (M+1)	

Example No.	304	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.21 (1H, s), 7.94 (1H, d, J=8.7Hz), 7.84 (1H, d, J=9.1Hz), 7.70 (1H, s), 7.26-7.39 (9H, m), 7.11 (2H, d, J=8.4Hz), 5.11 (2H, s), 4.26 (1H, m), 3.01 (3H, s), 2.97 (3H, s), 2.38-2.19 (2H, m), 1.97-1.78 (4H, m), 1.72-1.57 (1H, m), 1.48-1.17 (3H, m)
Purity	> 90% (NMR)	
MS	608 (M+1)	

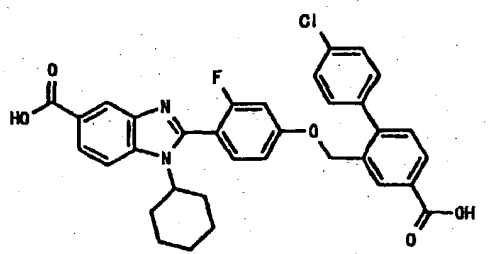
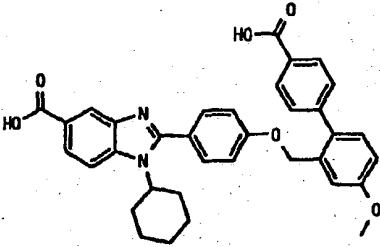
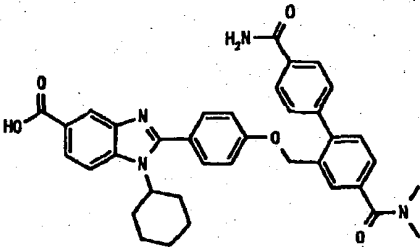
Example No.	305	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.24 (2H, s), 8.03 (1H, d, J=8.0Hz), 7.96 (1H, d, J=8.8Hz), 7.87 (1H, d, J=9.1Hz), 7.60-7.46 (6H, m), 7.09 (1H, dd, J=12.0, 1.8Hz), 6.97 (1H, dd, J=8.4, 1.8Hz), 5.16 (2H, s), 3.97 (1H, m), 2.31-2.11 (2H, m), 1.92-1.73 (4H, m), 1.70-1.57 (1H, m), 1.46-1.13 (3H, m)
Purity	> 90% (NMR)	
MS	599 (M+1)	

Table 204

Example No.	306	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.84 (1H, brs), 8.21 (1H, s), 7.98-7.84 (5H, m), 7.58 (2H, d, J=8.7Hz), 7.54 (2H, d, J= 7.8Hz), 7.34 (1H, d, J=8.7H z), 7.26 (1H, d, J=2.4Hz), 7. 13-7.06 (3H, m), 5.06 (2H, s) , 4.26 (1H, brt, J=12.7Hz), 3 .84 (3H, s), 2.36-2.17 (2H, b rm), 1.99-1.80 (4H, brm), 1. 73-1.59 (1H, brm), 1.47-1.1 7 (3H, brm)
Purity	> 90% (NMR)	
MS	577 (M+1)	

Example No.	307	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.22 (1H, s), 8.04 (1H, s), 7. 96 (2H, d, J=8.1Hz), 7.87 (2H , s), 7.72 (1H, d, J=1.2Hz), 7 .59-7.41 (7H, m), 5.12 (2H, s ), 4.25 (1H, brt, J=11.8Hz), 3.02 (3H, brs), 2.98 (3H, brs ), 2.38-2.15 (2H, brm), 1.93 -1.76 (4H, brm), 1.71-1.59 ( 1H, brm), 1.46-1.16 (3H, brm )
Purity	> 90% (NMR)	
MS	617 (M+1)	

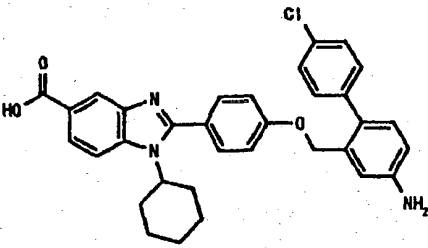
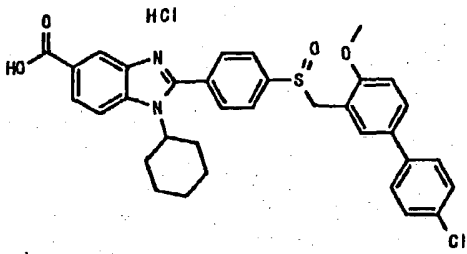
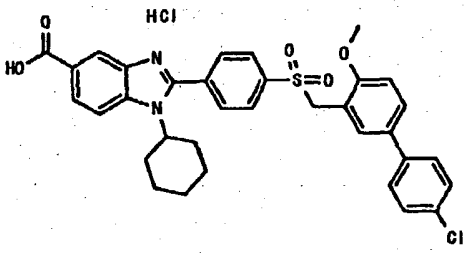
Example No.	308	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.27 (1H, s), 8.08 (1H, d, J=9 .0Hz), 7.93 (1H, d, J=8.7Hz) , 7.65 (2H, d, J=8.7Hz), 7.46 (2H, d, J=8.1Hz), 7.42 (2H, d , J=8.4Hz), 7.30-7.04 (5H, m ), 5.03 (2H, s), 4.32 (1H, m), 2.40-2.10 (2H, m), 2.05-1.1 0 (8H, m)
Purity	> 90% (NMR)	
MS	552 (M+1)	



Table 205

Example No.	309	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.15 and 7.99 (2H, ABq, J=8.9Hz), 7.84 and 7.59 (4H, A'B'q, J=8.3Hz), 7.46 (2H, d, J=8.4Hz), 7.22-7.16 (3H, m), 7.01-6.98 (2H, m), 4.27 and 4.23 (2H, A''B''q, J=12.9Hz), 3.78 (3H, s), 2.39-2.21 (2H, brm), 2.07-1.95 (2H, brm), 1.91-1.80 (2H, brm), 1.72-1.59 (1H, brm), 1.49-1.17 (3H, brm)
Purity	> 90% (NMR)	
MS		

Example No.	310	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.09 and 7.95 (2H, ABq, J=8.7Hz), 7.87 and 7.71 (4H, A'B'q, J=8.0Hz), 7.43 (2H, d, J=7.8Hz), 7.15 (1H, d, J=8.7Hz), 7.07-7.02 (4H, m), 4.66 (2H, s), 4.23 (1H, brt, J=11.8Hz), 3.76 (3H, s), 2.38-2.20 (2H, brm), 2.04-1.93 (2H, brm), 1.89-1.79 (2H, brm), 1.70-1.59 (1H, brm), 1.49-1.18 (3H, brm)
Purity	> 90% (NMR)	
MS	615 (M+1)	

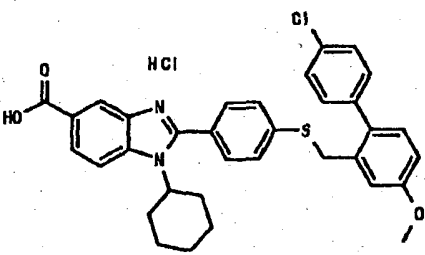
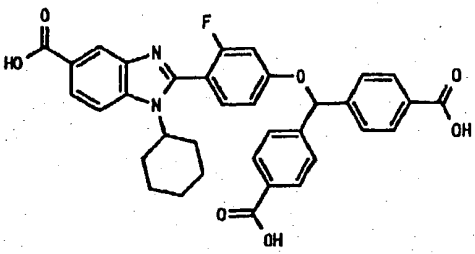
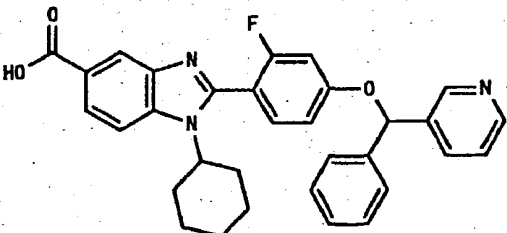
Example No.	311	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.21 and 8.01 (2H, ABq, J=8.7Hz), 7.65 (2H, d, J=8.4Hz), 7.52-7.41 (6H, m), 7.20 (1H, d, J=8.4Hz), 7.14 (1H, d, J=2.7Hz), 6.97 (1H, dd, J=8.4, 2.4Hz), 4.31 (1H, brt, J=9.8Hz), 4.28 (2H, s), 3.78 (3H, s), 2.37-2.20 (2H, brm), 2.07-1.95 (2H, brm), 1.92-1.80 (2H, brm), 1.71-1.60 (1H, brm), 1.50-1.19 (3H, brm)
Purity	> 90% (NMR)	
MS	583 (M+1)	

Table 206

Example No.	312	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.22(1H, s), 8.12(1H, d, J=8.4Hz), 8.00-7.84(5H, m), 7.70(4H, d, J=8.4Hz), 7.56(1H, t, J=8.6Hz), 7.23(1H, d, J=12.0Hz), 7.13(1H, d, J=8.6Hz), 6.97(1H, s), 3.92(1H, m), 2.35-2.00(2H, m), 1.95-1.70(4H, m), 1.65-1.55(1H, m), 1.50-1.05(3H, m)
Purity	> 90% (NMR)	
MS	609 (M+1)	

Example No.	313	1H NMR(δ) ppm
		300MHz, DMSO-d6 8.89(1H, brs), 8.63(1H, brs), 8.24(1H, s), 8.11(1H, d, J=7.8Hz), 7.99(1H, d, J=8.8Hz), 7.89(1H, d, J=9.9Hz), 7.61-7.55(4H, m), 7.43(2H, t, J=7.7Hz), 7.34(1H, t, J=7.2Hz), 7.24(1H, d, J=12.0Hz), 7.14(1H, d, J=8.6Hz), 6.95(1H, s), 3.96(1H, m), 2.35-2.05(2H, m), 2.00-1.50(5H, m), 1.45-1.10(3H, m)
Purity	> 90% (NMR)	
MS	522 (M+1)	

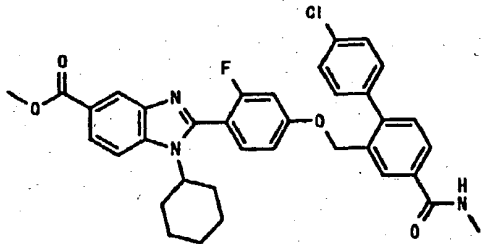
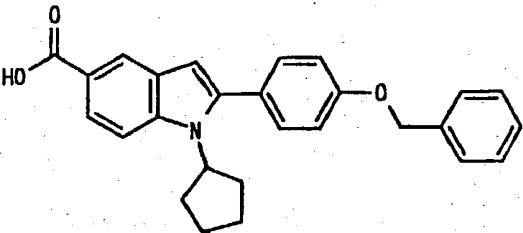
Example No.	314	1H NMR(δ) ppm
		300MHz, CDCl3 8.48(1H, d, J=1.4Hz), 8.05(1H, d, J=1.8Hz), 8.98(1H, d, J=8.6Hz), 7.82(1H, d, J=7.9Hz), 7.66(1H, d, J=8.6Hz), 7.55-7.24(6H, m), 6.78(1H, d, J=8.6), 2.69(1H, d, J=11.6Hz), 2.2Hz), 6.40-6.30(1H, m), 4.99(2H, s), 4.02(1H, m), 3.95(3H, s), 3.05(3H, d, J=4.8Hz), 2.32-2.13(2H, m), 2.03-1.87(4H, m), 1.81-1.71(1H, m), 1.46-1.23(3H, m)
Purity	> 90% (NMR)	
MS	626 (M+1)	

Table 207

Example No.	503	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.23 (1H, s), 7.76 (1H, d, J=8.7Hz), 7.58 (1H, d, J=8.8Hz), 7.51-7.32 (7H, m), 7.17 (2H, d, J=8.7Hz), 6.55 (1H, s), 5.18 (2H, s), 4.75 (1H, m), 2.35-2.12 (2H, m), 2.10-1.85 (4H, m), 1.80-1.50 (2H, m)
Purity	> 90% (NMR)	
MS	412 (M+1)	

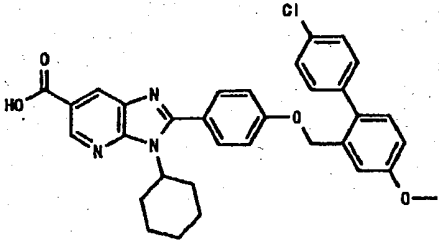
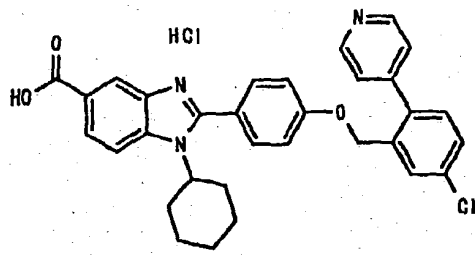
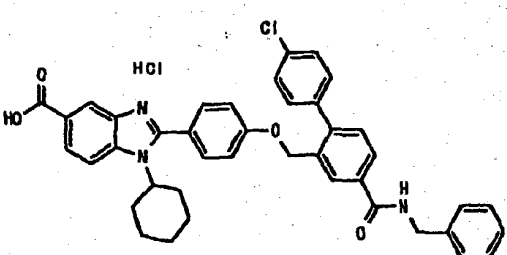
Example No.	701	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.96 (1H, s), 8.50 (1H, s), 7.77 (2H, d, J=8.7Hz), 7.50-7.40 (4H, m), 7.30 (1H, d, J=8.4Hz), 7.24 (1H, d, J=2.4Hz), 7.16 (2H, d, J=8.4Hz), 7.06 (1H, dd, J=2.4Hz, 8.1Hz), 5.06 (2H, s), 4.31 (1H, s), 3.83 (3H, s), 2.80-2.55 (2H, m), 2.00-1.80 (4H, m), 1.70-1.55 (1H, m), 1.40-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	568 (M+1)	

Table 208

Example No.	315	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.84 (2H, d, J=6.3Hz), 8.28 (1H, s), 8.17 and 7.99 (2H, ABq, J=8.7Hz), 7.87-7.85 (3H, m), 7.70-7.50 (3H, m), 7.52 (1H, d, J=8.3Hz), 7.18 (2H, d, J=8.7Hz), 5.22 (2H, s), 4.31 (1H, br t, J=12.5Hz), 2.36-2.18 (2H, m), 2.03-1.78 (4H, m), 1.70-1.58 (1H, m), 1.50-1.23 (3H, m)
Purity	> 90% (NMR)	
MS	538 (M+1)	

Example No.	316	1H NMR (δ) ppm
		300MHz, DMSO-d6 9.23 (1H, t, J=6.3Hz), 8.29 (1H, s), 8.25-8.22 (2H, m), 8.03 (2H, d, J=7.9Hz), 7.55-7.48 (5H, m), 7.34 (4H, d, J=4.4Hz), 7.28-7.22 (3H, m), 5.15 (2H, s), 4.52 (2H, d, J=5.9Hz), 4.35 (1H, br t, J=12.1Hz), 2.37-2.18 (2H, m), 2.08-1.95 (2H, m), 1.91-1.79 (2H, m), 1.72-1.59 (1H, m), 1.47-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	670 (M+1)	

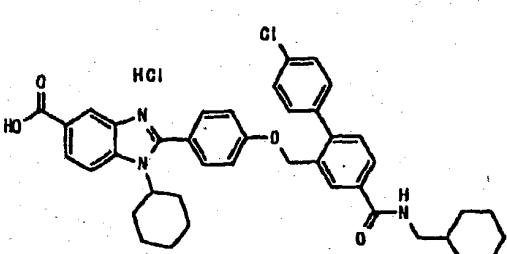
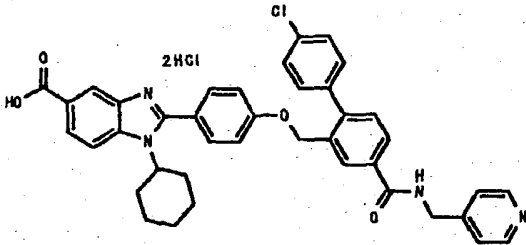
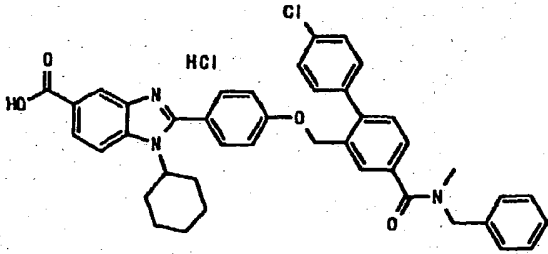
Example No.	317	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.59 (1H, t, J=5.5Hz), 8.28 (1H, s), 8.21 and 8.01 (2H, ABq, J=8.8Hz), 8.16 (1H, s), 7.97 and 7.46 (2H, A'B'q, J=8.0Hz), 7.71 and 7.23 (4H, A''B''q, J=8.7Hz), 7.53 and 7.49 (4H, A'B''q, J=9.2Hz), 5.14 (2H, s), 4.34 (1H, br t, J=12.8Hz), 3.14 (2H, t, J=6.3Hz), 2.38-2.18 (2H, m), 2.07-1.78 (4H, m), 1.78-1.47 (7H, m), 1.47-1.07 (6H, m), 1.03-0.83 (2H, m)
Purity	> 90% (NMR)	
MS	676 (M+1)	

Table 209

Example No.	318	1H NMR (δ) ppm
		300MHz, DMSO-d6 9.63 (1H, t, J=4.8Hz), 8.86 and 7.97 (4H, ABq, J=6.6Hz), 8.30 (1H, s), 8.27 (1H, s), 8.23 and 8.03 (2H, A 'B' q, J=8.8Hz), 8.09 and 7.54 (2 H, A''B'' q, J=8.1Hz), 7.73 and 7.2 4 (4H, A'''B''' q, J=8.8Hz), 7.54a nd 7.52 (4H, A''''B'''' q, J=8.8Hz), 5.16 (2H, s), 4.78 (2H, d, J=5.6Hz ), 4.35 (1H, br t, J=11.0Hz), 2.39-2.19 (2H, m) , 2.07-1.96 (2H, m), 1.91-1.78 (2H, m), 1.70-1.57 (1H, m) 1.50-1.19 (3H, m).
Purity	> 90% (NMR)	
MS	671 (M+1)	

Example No.	319	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.28 (1H, s), 8.24 and 8.03 (2H, A Bq, J=9.0Hz), 7.77 (1H, s), 7.70 (2H, d, J=8.4Hz), 7.64-7.10 (13 H, m), 5.16 (2H, s), 4.74 and 4.57 (total 2H, each br s), 4.34 (1H, br t, J=11.7Hz), 2.90 (3H, s), 2.35 -2.17 (2H, m), 2.07-1.93 (2H, m) , 1.93-1.78 (2H, m), 1.71-1.57 (1H, m), 1.51-1.19 (3H, m)
Purity	> 90% (NMR)	
MS	684 (M+1)	

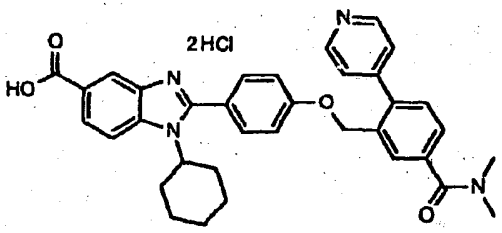
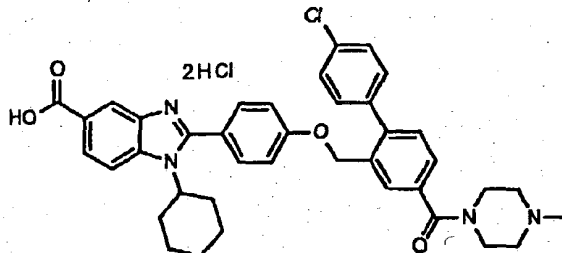
Example No.	320	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.94 and 8.06 (4H, ABq, J=6.8Hz) , 8.33 (1H, s), 8.28 and 8.05 (2H, A'B' q, J=8.7Hz), 7.80 (1H, s), 7 .73 and 7.22 (4H, A''B'' q, J=8.7Hz ), 7.63 and 7.57 (2H, A'''B''' q, J= 7.9Hz), 5.30 (2H, s), 4.34 (1H, b r t, J=12.1Hz), 3.04 (3H, s), 2.97 (3H, s), 2.38-2.18 (2H, m), 2.10 -1.96 (2H, m), 1.93-1.80 (2H, m) , 1.72-1.58 (1H, m), 1.52-1.08 (3H, m)
Purity	> 90% (NMR)	
MS	575 (M+1)	

Table 210

Example No.	321	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 11.19 (1H, br s), 8.31 (1H, s), 8.23 and 8.02 (2H, ABq, J=9.0Hz), 7.77 (1H, s), 7.72 and 7.23 (4H, A'B'q, J=8.7Hz), 7.59 and 7.48 (2H, A''B''q, J=7.9Hz), 7.53 and 7.51 (4H, A'''B'''q, J=9.0Hz), 5.16 (2H, s), 4.72-2.97 (8H, br m), 4.34 (1H, br t, J=12.1Hz), 2.79 (3H, s), 2.38-2.17 (2H, m), 2.07-1.93 (2H, m), 1.93-1.78 (2H, m), 1.69-1.58 (1H, m), 1.50-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	663 (M+1)	

Example No.	322	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9.54 (1H, t, J=5.7Hz), 8.91 (1H, s), 8.81 (1H, d, J=4.9Hz), 8.48 (1H, d, J=7.9Hz), 8.32 (1H, s), 8.27 (1H, d, J=9.0Hz), 8.25 (1H, s), 8.07-7.97 (3H, m), 7.74 and 7.25 (4H, ABq, J=8.9Hz), 7.56-7.49 (5H, m), 5.16 (2H, s), 4.69 (2H, d, J=5.6Hz), 4.36 (1H, br t, J=12.4Hz), 2.37-2.20 (2H, m), 2.09-1.97 (2H, m), 1.91-1.78 (2H, m), 1.70-1.57 (1H, m), 1.50-1.17 (3H, m)
Purity	> 90% (NMR)	
MS	671 (M+1)	

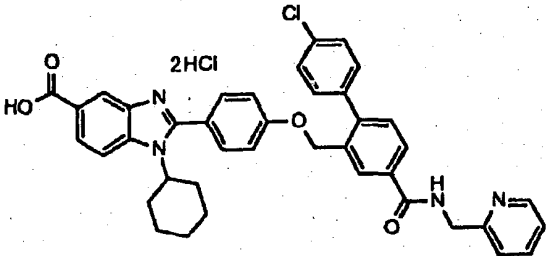
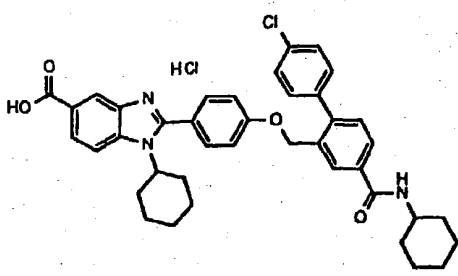
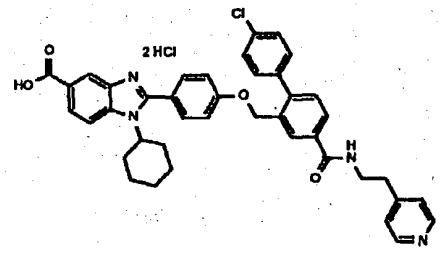
Example No.	323	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9.52 (1H, t, J=6.0Hz), 8.72 (1H, d, J=5.3Hz), 8.30-8.19 (4H, m), 8.08 (1H, d, J=7.9Hz), 8.02 (1H, d, J=7.6Hz), 7.77-7.64 (4H, m), 7.57-7.49 (5H, m), 7.24 (2H, d, J=8.7Hz), 5.16 (2H, s), 4.77 (2H, d, J=5.6Hz), 4.34 (1H, t, J=12.8Hz), 2.36-2.19 (2H, m), 2.07-1.95 (2H, m), 1.91-1.78 (2H, m), 1.69-1.59 (1H, m), 1.45-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	671 (M+1)	

Table 211

Example No.	324	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.36 (1H, d, J=7.9Hz), 8.30 (1H, s), 8.28 and 8.05 (2H, ABq, J=8.8 Hz), 8.16 (1H, s), 7.79 and 7.46 (2H, A'B'q, J=8.3Hz), 7.74 and 7.25 (4H, A''B''q, J=8.9Hz), 7.52 and 7.50 (4H, A'''B'''q, J=8.7Hz), 5.14 (2H, s), 4.36 (1H, br t, J=12.1Hz), 3.80 (1H, br s), 2.39-2.18 (2H, m), 2.10-1.98 (2H, m), 1.93-1.57 (8H, m), 1.49-1.04 (8H, m)
Purity	> 90% (NMR)	
MS	662 (M+1)	

Example No.	325	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.86 (1H, t, J=6.0Hz), 8.84 and 8.00 (4H, ABq, J=6.6Hz), 8.33 (1H, s), 8.27 and 8.04 (2H, A'B'q, J=9.0Hz), 8.12 (1H, s), 7.92 and 7.46 (2H, A''B''q, J=7.9Hz), 7.74 and 7.23 (4H, A'''B'''q, J=9.0Hz), 7.53 and 7.49 (4H, A''''B''''q, J=9.1 Hz), 5.13 (2H, s), 4.36 (1H, br t, J=12.8Hz), 3.70 (2H, td, J=6.8, 6.0Hz), 3.21 (2H, t, J=6.8Hz), 2.38-2.20 (2H, m), 2.09-1.95 (2H, m), 1.91-1.77 (2H, m), 1.70-1.59 (1H, m), 1.49-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	685 (M+1)	

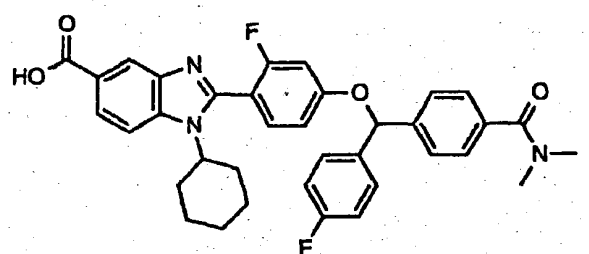
Example No.	326	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 12.80 (1H, brs), 8.23 (1H, s), 7.90 (1H, d, J=8.7Hz), 7.83 (1H, d, J=8.7Hz), 7.60-7.50 (5H, m), 7.39 (2H, d, J=7.8Hz), 7.23-7.10 (3H, m), 7.05 (1H, d, J=7.8Hz), 6.85 (1H, s), 3.94 (1H, s), 2.97, 2.88 (6H, s), 2.30-2.10 (2H, m), 1.90-1.50 (5H, m), 1.40-1.00 (3H, m)
Purity	> 90% (NMR)	
MS	610 (M+1)	

Table 212

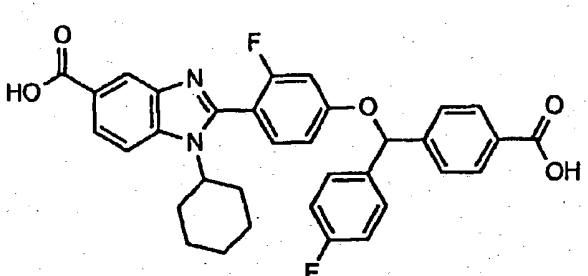
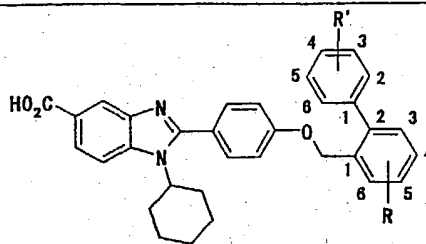
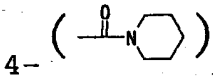
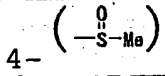
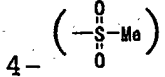
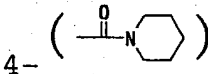
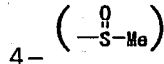
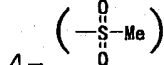
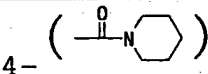
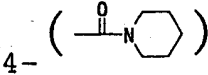
Example No.	327	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.20-12.60 (2H, brs), 8.23 (1H, s), 7.98 (2H, d, J=6.6Hz), 7.95 (1H, d, J=8.7Hz), 7.87 (1H, d, J=8.7Hz), 7.70-7.50 (5H, m), 7.27-7.20 (3H, m), 7.08 (1H, d, J=7.8Hz), 6.90 (1H, s), 3.93 (1H, s), 2.51-2.05 (2H, m), 1.90-1.70 (4H, m), 1.65-1.55 (1H, m), 1.40-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	583 (M+1)	

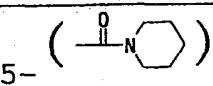
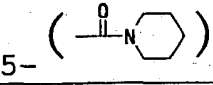
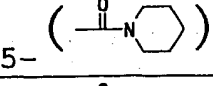
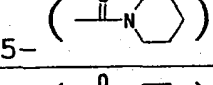
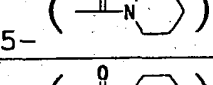
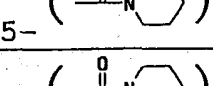
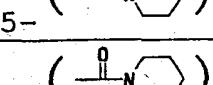

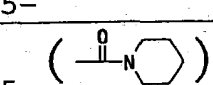
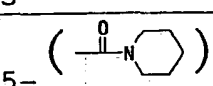
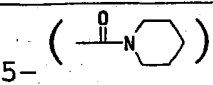
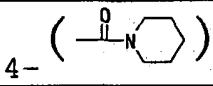
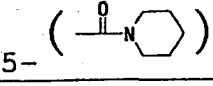
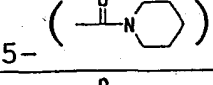
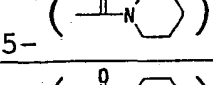
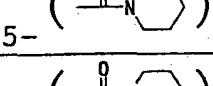
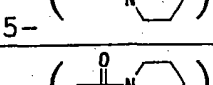
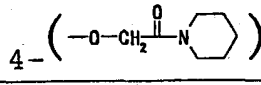
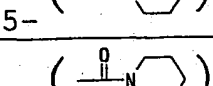
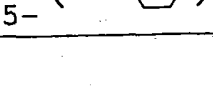

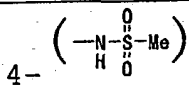


Table 213



Ex. No.	R	R'
2001	-H	4-(-Me)
2002	-H	3-(-CF <sub>3</sub> )
2003	5-(-F)	-H
2004	3-(-F)	2-(-F)
2005	3-(-F)	3-(-F)
2006	3-(-F)	4-(-F)
2007	4-(-F)	4-(-F)
2008	5-(-F)	4-(-F)
2009	6-(-F)	4-(-F)
2010	4-(-F)	4-(-Cl)
2011	5-(-F)	4-(-Me)
2012	5-(-F)	4-(-CF <sub>3</sub> )
2013	5-(-F)	4-(-CO <sub>2</sub> H)
2014	5-(-F)	4-(-CO <sub>2</sub> Me)
2015	5-(-F)	4- 
2016	5-(-F)	4-(-CONH <sub>2</sub> )
2017	5-(-F)	4-{-CON(Me) <sub>2</sub> }
2018	5-(-F)	4-(-OMe)
2019	5-(-F)	4-(-SMe)
2020	5-(-F)	4- 
2021	5-(-F)	4- 
2022	4-(-Cl)	-H

2023	4-(-Cl)	4-(-F)
2024	4-(-Cl)	4-(-Cl)
2025	4-(-Cl)	4-(-Me)
2026	5-(-Cl)	4-(-CF <sub>3</sub> )
2027	4-(-Cl)	4-(-CO <sub>2</sub> H)
2028	5-(-Cl)	4-(-CO <sub>2</sub> Me)
2029	5-(-Cl)	4- 
2030	4-(-Cl)	4-(-CONH <sub>2</sub> )
2031	5-(-Cl)	4-(-CON(Me) <sub>2</sub> )
2032	5-(-Cl)	3-(-OMe)
2033	4-(-Cl)	4-(-SMe)
2034	5-(-Cl)	4- 
2035	4-(-Cl)	4- 
2036	5-(-CN)	4-(-F)
2037	4-(-CN)	4-(-Cl)
2038	5-(-NO <sub>2</sub> )	4-(-F)
2039	4-(-NO <sub>2</sub> )	4-(-Cl)
2040	5-(-Me)	4-(-CO <sub>2</sub> H)
2041	5-(-Me)	4-(-CO <sub>2</sub> Me)
2042	5-(-Me)	4- 
2043	5-(-CF <sub>3</sub> )	4-(-CO <sub>2</sub> H)
2044	5-(-CF <sub>3</sub> )	4-(-CO <sub>2</sub> Me)
2045	5-(-CF <sub>3</sub> )	4- 
2046	5-(-CO <sub>2</sub> H)	4-(-F)
2047	4-(-CO <sub>2</sub> H)	4-(-Cl)
2048	5-(-CO <sub>2</sub> Me)	4-(-F)
2049	5-(-CO <sub>2</sub> Me)	4-(-Cl)
2050	5-(-Ac)	4-(-F)

2051	5-(-Ac)	4-(-Cl)
2052	5- 	-H
2053	5- 	4-(-F)
2054	5- 	4-(-Cl)
2055	5- 	4-(-CN)
2056	5- 	4-(-NO <sub>2</sub> )
2057	5- 	4-(-Me)
2058	5- 	4-(-CF <sub>3</sub> )
2059	5- 	4-(-Ac)
2060	5- 	4-(-CO <sub>2</sub> H)
2061	5- 	4-(-CO <sub>2</sub> Me)
2062	5- 	4- 
2063	5- 	4-(-CONH <sub>2</sub> )
2064	5- 	4-{-CON(Me) <sub>2</sub> }
2065	5- 	4-{-C(=NH)NH <sub>2</sub> }
2066	5- 	4-(-OMe)
2067	5- 	4- 
2068	5- 	4-(-NHMe)
2069	5- 	4-(-NHAc)
2070	5- 	4- 

2071	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- (-SMe)
2072	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{Me} \right)$
2073	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{Me} \right)$
2074	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{NH}_2 \right)$
2075	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left\{ \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{N}(\text{Me})_2 \right\}$
2076	5- (-CONH <sub>2</sub> )	-H
2077	5- (-CONH <sub>2</sub> )	4- (-F)
2078	5- (-CONH <sub>2</sub> )	2,3,4,5,6-penta- (-F)
2079	5- (-CONH <sub>2</sub> )	2- (-Cl)
2080	5- (-CONH <sub>2</sub> )	3- (-Cl)
2081	3- (-CONH <sub>2</sub> )	2- (-Cl)
2082	3- (-CONH <sub>2</sub> )	3- (-Cl)
2083	3- (-CONH <sub>2</sub> )	4- (-Cl)
2084	4- (-CONH <sub>2</sub> )	2- (-Cl)
2085	4- (-CONH <sub>2</sub> )	3- (-Cl)
2086	4- (-CONH <sub>2</sub> )	4- (-Cl)
2087	6- (-CONH <sub>2</sub> )	2- (-Cl)
2088	6- (-CONH <sub>2</sub> )	3- (-Cl)
2089	6- (-CONH <sub>2</sub> )	4- (-Cl)
2090	5- (-CONH <sub>2</sub> )	3,5-di- (-Cl)
2091	5- (-CONH <sub>2</sub> )	4- (-CN)
2092	5- (-CONH <sub>2</sub> )	4- (-NO <sub>2</sub> )
2093	5- (-CONH <sub>2</sub> )	4- (-Me)
2094	5- (-CONH <sub>2</sub> )	2,6-di- (-Me)
2095	5- (-CONH <sub>2</sub> )	4- (-CF <sub>3</sub> )
2096	5- (-CONH <sub>2</sub> )	4- (-Ac)
2097	5- (-CONH <sub>2</sub> )	4- (-CO <sub>2</sub> H)
2098	5- (-CONH <sub>2</sub> )	4- (-CO <sub>2</sub> Me)

2099	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array} \text{C}_6\text{H}_{11} \right)$
2100	5-(-CONH <sub>2</sub> )	4-(-CONH <sub>2</sub> )
2101	5-(-CONH <sub>2</sub> )	3,5-di-(-CONH <sub>2</sub> )
2102	5-(-CONH <sub>2</sub> )	4-{-CON(Me) <sub>2</sub> }
2103	5-(-CONH <sub>2</sub> )	4-{-C(=NH)NH <sub>2</sub> }
2104	5-(-CONH <sub>2</sub> )	4-(-OMe)
2105	5-(-CONH <sub>2</sub> )	3,4,5-tri-(-OMe)
2106	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N} \end{array} \text{C}_6\text{H}_{11} \right)$
2107	5-(-CONH <sub>2</sub> )	4-(-NHMe)
2108	5-(-CONH <sub>2</sub> )	4-(-NHAc)
2109	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\ \mid \\ \text{H} \end{array} \right)$
2110	5-(-CONH <sub>2</sub> )	4-(-SMe)
2111	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2112	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \mid \\ \text{O} \end{array} \right)$
2113	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \mid \\ \text{O} \end{array} \right)$
2114	5-(-CONH <sub>2</sub> )	4- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array} \right\}$
2115	5-{-CON(Me) <sub>2</sub> }	-H
2116	5-{-CON(Me) <sub>2</sub> }	4-(-F)
2117	4-{-CON(Me) <sub>2</sub> }	4-(-Cl)
2118	5-{-CON(Me) <sub>2</sub> }	4-(-CN)
2119	5-{-CON(Me) <sub>2</sub> }	4-(-NO <sub>2</sub> )
2120	5-{-CON(Me) <sub>2</sub> }	4-(-Me)
2121	4-{-CON(Me) <sub>2</sub> }	4-(-CF <sub>3</sub> )
2122	5-{-CON(Me) <sub>2</sub> }	4-(-Ac)
2123	5-{-CON(Me) <sub>2</sub> }	4-(-CO <sub>2</sub> H)
2124	5-{-CON(Me) <sub>2</sub> }	4-(-CO <sub>2</sub> Me)

2125	5-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{N} \end{array} \begin{array}{c} \diagup \\ \diagdown \end{array} \right)$
2126	5-{-CON(Me) <sub>2</sub> }	3-{-CONH <sub>2</sub> }
2127	4-{-CON(Me) <sub>2</sub> }	4-{-CON(Me) <sub>2</sub> }
2128	5-{-CON(Me) <sub>2</sub> }	4-{-C(=NH)NH <sub>2</sub> }
2129	5-{-CON(Me) <sub>2</sub> }	4-{-OMe}
2130	5-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N} \end{array} \begin{array}{c} \diagup \\ \diagdown \end{array} \right)$
2131	5-{-CON(Me) <sub>2</sub> }	4-{-NHMe}
2132	5-{-CON(Me) <sub>2</sub> }	4-{-NHAc}
2133	5-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \parallel \\ \text{O} \end{array} \right)$
2134	4-{-CON(Me) <sub>2</sub> }	4-{-SMe}
2135	5-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$
2136	4-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$
2137	5-{-CON(Me) <sub>2</sub> }	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$
2138	5-{-CON(Me) <sub>2</sub> }	4- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$
2139	5-{-OMe}	-H
2140	5-{-OMe}	4-{-F}
2141	3-{-OMe}	4-{-Cl}
2142	4-{-OMe}	4-{-Cl}
2143	5-{-OMe}	2-{-Cl}
2144	5-{-OMe}	3-{-Cl}
2145	6-{-OMe}	4-{-Cl}
2146	5-{-OMe}	4-{-CN}
2147	5-{-OMe}	4-{-NO <sub>2</sub> }
2148	5-{-OMe}	4-{-Me}
2149	5-{-OMe}	4-{-CF <sub>3</sub> }
2150	5-{-OMe}	4-{-Ac}

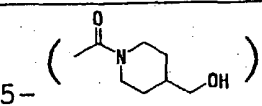
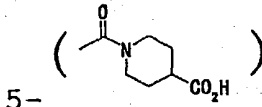
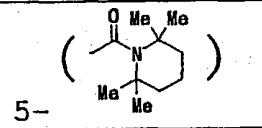
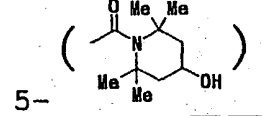
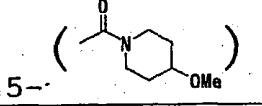
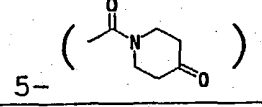
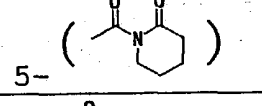
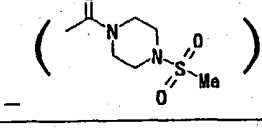
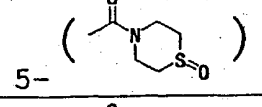
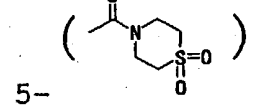
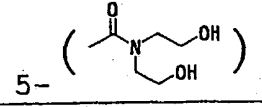
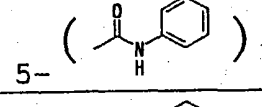
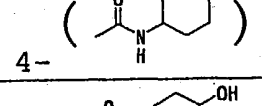
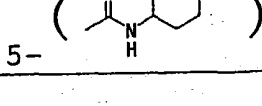
2151	4-(-OMe)	4-(-CO <sub>2</sub> H)
2152	4,5-di-(-OMe)	4-(-CO <sub>2</sub> H)
2153	5-(-OMe)	4-(-CO <sub>2</sub> Me)
2154	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array} \text{C}_6\text{H}_{11} \right)$
2155	5-(-OMe)	4-(-CONH <sub>2</sub> )
2156	5-(-OMe)	4-{-CON(Me) <sub>2</sub> }
2157	5-(-OMe)	4-{-C(=NH)NH <sub>2</sub> }
2158	5-(-OMe)	4-(-OMe)
2159	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N} \end{array} \text{C}_6\text{H}_{11} \right)$
2160	5-(-OMe)	4-(-NHMe)
2161	5-(-OMe)	4-(-NHAc)
2162	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \end{array} \right)$
2163	5-(-OMe)	4-(-SMe)
2164	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2165	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array} \right)$
2166	5-(-OMe)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array} \right)$
2167	5-(-OMe)	4- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array} \right\}$
2168	5-(-NHMe)	4-(-F)
2169	5-(-NHMe)	4-(-Cl)
2170	5-(-NHAc)	4-(-F)
2171	5-(-NHAc)	4-(-Cl)
2172	5-(-NHAc)	4-(-Ac)
2173	5-(-NHAc)	4-(-CONH <sub>2</sub> )
2174	5-(-NHAc)	4-{-CON(Me) <sub>2</sub> }
2175	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \end{array} \right)$	4-(-F)

2176	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-Cl)
2177	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-Me)
2178	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-CF <sub>3</sub> )
2179	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-CO <sub>2</sub> H)
2180	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-CO <sub>2</sub> Me)
2181	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---} \text{N} \text{---} \text{C}_6\text{H}_{11} \end{array} \right)$
2182	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- (-SMe)
2183	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2184	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array} \right)$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2185	5- (-SMe)	4- (-F)
2186	4- (-SMe)	4- (-Cl)
2187	5- (-SMe)	4- (-Me)
2188	5- (-SMe)	4- (-CF <sub>3</sub> )
2189	5- (-SMe)	4- (-Ac)
2190	5- (-SMe)	4- (-CONH <sub>2</sub> )
2191	5- (-SMe)	4- {-CON (Me) <sub>2</sub> }
2192	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-F)
2193	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Cl)
2194	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Me)
2195	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-CF <sub>3</sub> )
2196	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Ac)
2197	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-CONH <sub>2</sub> )



2198	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4-{-CON(Me) <sub>2</sub> }
2199	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-F)
2200	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-Cl)
2201	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-Me)
2202	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-CF <sub>3</sub> )
2203	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-Ac)
2204	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-(-CONH <sub>2</sub> )
2205	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4-{-CON(Me) <sub>2</sub> }
2206	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-F)
2207	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-Cl)
2208	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	2,4-di-(-Cl)
2209	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-Me)
2210	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	3-(-CF <sub>3</sub> )
2211	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-CF <sub>3</sub> )
2212	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-CONH <sub>2</sub> )
2213	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-{-CON(Me) <sub>2</sub> }
2214	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4-(-SMe)
2215	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2216	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$

2217	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-F)
2218	4- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-Cl)
2219	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-Me)
2220	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-CF <sub>3</sub> )
2221	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-CONH <sub>2</sub> )
2222	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- {-CON (Me) <sub>2</sub> }
2223	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- (-SMe)
2224	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$
2225	5- $\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$
2226	5- {-O- (CH <sub>2</sub> ) <sub>2</sub> -OH}	4- (-Cl)
2227	5- {-O- (CH <sub>2</sub> ) <sub>3</sub> -OH}	4- (-Cl)
2228	5- $\left( \text{---O---} \begin{array}{c} \text{Cyclohexyl} \end{array} \right)$	4- (-Cl)
2229	5- $\left( \text{---O---} \begin{array}{c} \text{Pyridyl} \end{array} \right)$	4- (-Cl)
2230	5- $\left( \text{---O---} \begin{array}{c} \text{2-Methylthiazolyl} \end{array} \right)$	4- (-Cl)
2231	5- $\left( \text{---O---} \begin{array}{c} \text{4-Hydroxypiperidinyl} \end{array} \right)$	4- (-Cl)
2232	5- $\left( \text{---O---} \begin{array}{c} \text{N-C(=O)-CH}_2\text{---} \\ \text{4-Hydroxypiperidinyl} \end{array} \right)$	4- (-Cl)
2233	5- $\left( \begin{array}{c} \text{N-C(=O)-CH}_3 \\ \text{4-Hydroxypyrrolidinyl} \end{array} \right)$	4- (-Cl)
2234	5- $\left( \begin{array}{c} \text{N-C(=O)-CH}_3 \\ \text{4-Hydroxypiperidinyl} \end{array} \right)$	4- (-Cl)
2235	5- $\left( \begin{array}{c} \text{N-C(=O)-CH}_3 \\ \text{4,6-Dihydroxypiperidinyl} \end{array} \right)$	4- (-Cl)

2236	5- 	4- (-Cl)
2237	5- 	4- (-Cl)
2238	5- 	4- (-Cl)
2239	5- 	4- (-Cl)
2240	5- 	4- (-Cl)
2241	5- 	4- (-Cl)
2242	5- 	4- (-Cl)
2243	5- 	4- (-Cl)
2244	5- 	4- (-Cl)
2245	5- 	4- (-Cl)
2246	5- 	4- (-Cl)
2247	5- 	4- (-Cl)
2248	4- 	4- (-Cl)
2249	5- 	4- (-Cl)

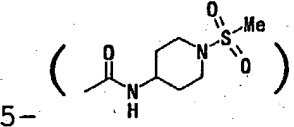
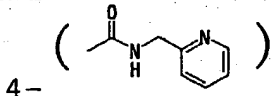
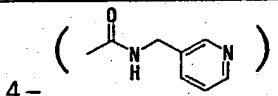
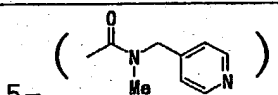
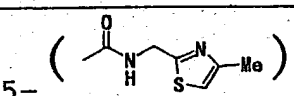
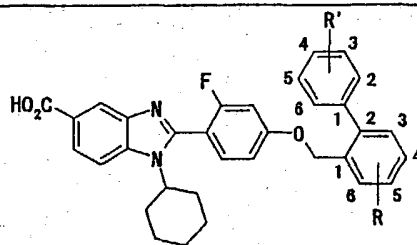
2250	5- 	4-(-Cl)
2251	4- 	4-(-Cl)
2252	4- 	4-(-Cl)
2253	5- 	4-(-Cl)
2254	5- 	4-(-Cl)

Table 214



Ex. No.	R	R'
2255	-H	-H
2256	-H	4-(-Me)
2257	-H	3-(-CF <sub>3</sub> )
2258	5-(-F)	-H
2259	5-(-F)	4-(-F)
2260	5-(-F)	4-(-Cl)
2261	5-(-F)	4-(-Me)
2262	5-(-F)	4-(-CF <sub>3</sub> )
2263	5-(-F)	4-(-CO <sub>2</sub> H)
2264	5-(-F)	4-(-CO <sub>2</sub> Me)
2265	5-(-F)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{Cyclohexyl} \right)$
2266	5-(-F)	4-(-CONH <sub>2</sub> )
2267	5-(-F)	4-(-CON(Me) <sub>2</sub> )
2268	5-(-F)	4-(-OMe)
2269	5-(-F)	4-(-SMe)
2270	5-(-F)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---S---} \end{array} \text{Me} \right)$
2271	5-(-F)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---S---} \\ \parallel \\ \text{O} \end{array} \text{Me} \right)$
2272	4-(-Cl)	-H
2273	5-(-Cl)	4-(-F)
2274	4-(-Cl)	4-(-Cl)
2275	5-(-Cl)	4-(-Me)
2276	5-(-Cl)	4-(-CF <sub>3</sub> )

2277	5-(-Cl)	4-(-CO <sub>2</sub> H)
2278	5-(-Cl)	4-(-CO <sub>2</sub> Me)
2279	5-(-Cl)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{C}_6\text{H}_{11} \right)$
2280	5-(-Cl)	4-(-CONH <sub>2</sub> )
2281	5-(-Cl)	4-{-CON(Me) <sub>2</sub> }
2282	5-(-Cl)	4-(-OMe)
2283	5-(-Cl)	4-(-SMe)
2284	5-(-Cl)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---S---} \end{array} \text{Me} \right)$
2285	5-(-Cl)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---S---} \\ \parallel \\ \text{O} \end{array} \text{Me} \right)$
2286	5-(-CN)	4-(-F)
2287	5-(-CN)	4-(-Cl)
2288	5-(-NO <sub>2</sub> )	4-(-F)
2289	5-(-NO <sub>2</sub> )	4-(-Cl)
2290	5-(-Me)	4-(-CO <sub>2</sub> H)
2291	5-(-Me)	4-(-CO <sub>2</sub> Me)
2292	5-(-Me)	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{C}_6\text{H}_{11} \right)$
2293	5-(-CF <sub>3</sub> )	4-(-CO <sub>2</sub> H)
2294	5-(-CF <sub>3</sub> )	4-(-CO <sub>2</sub> Me)
2295	5-(-CF <sub>3</sub> )	4- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{C}_6\text{H}_{11} \right)$
2296	5-(-CO <sub>2</sub> H)	4-(-F)
2297	4-(-CO <sub>2</sub> H)	4-(-Cl)
2298	5-(-CO <sub>2</sub> Me)	4-(-F)
2299	5-(-CO <sub>2</sub> Me)	4-(-Cl)
2300	5-(-Ac)	4-(-F)
2301	5-(-Ac)	4-(-Cl)
2302	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{C}_6\text{H}_{11} \right)$	-H
2303	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ \text{---N---} \end{array} \text{C}_6\text{H}_{11} \right)$	4-(-F)


2304	$4-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---Cl})$
2305	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---CN})$
2306	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---NO}_2)$
2307	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---Me})$
2308	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---CF}_3)$
2309	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---Ac})$
2310	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---CO}_2\text{H})$
2311	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---CO}_2\text{Me})$
2312	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$
2313	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---CONH}_2)$
2314	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}\{\text{---CON}(\text{Me})_2\}$
2315	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}\{\text{---C}(=\text{NH})\text{NH}_2\}$
2316	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---OMe})$
2317	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\left(\text{---O---CH}_2\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$
2318	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---NHMe})$
2319	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---NHAc})$
2320	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\overset{\text{O}}{\parallel}\text{S}\text{---Me}\right)$
2321	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\text{---}(\text{---SMe})$
2322	$5-\left(\text{---}\overset{\text{O}}{\parallel}\text{N}\text{---}\right)$	$4-\left(\text{---}\overset{\text{O}}{\parallel}\text{S}\text{---Me}\right)$

2323	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{Me} \right)$
2324	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{NH}_2 \right)$
2325	5- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$	4- $\left\{ \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{N}(\text{Me})_2 \right\}$
2326	5- $(-\text{CONH}_2)$	-H
2327	5- $(-\text{CONH}_2)$	4- $(-\text{F})$
2328	4- $(-\text{CONH}_2)$	4- $(-\text{Cl})$
2329	5- $(-\text{CONH}_2)$	4- $(-\text{CN})$
2330	5- $(-\text{CONH}_2)$	4- $(-\text{NO}_2)$
2331	5- $(-\text{CONH}_2)$	4- $(-\text{Me})$
2332	5- $(-\text{CONH}_2)$	4- $(-\text{CF}_3)$
2333	5- $(-\text{CONH}_2)$	4- $(-\text{Ac})$
2334	5- $(-\text{CONH}_2)$	4- $(-\text{CO}_2\text{H})$
2335	5- $(-\text{CONH}_2)$	4- $(-\text{CO}_2\text{Me})$
2336	5- $(-\text{CONH}_2)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$
2337	5- $(-\text{CONH}_2)$	4- $(-\text{CONH}_2)$
2338	5- $(-\text{CONH}_2)$	4- $\{ -\text{CON}(\text{Me})_2 \}$
2339	5- $(-\text{CONH}_2)$	4- $\{ -\text{C}(=\text{NH})\text{NH}_2 \}$
2340	5- $(-\text{CONH}_2)$	4- $(-\text{OMe})$
2341	5- $(-\text{CONH}_2)$	4- $\left( \text{---} \text{O} \text{---} \text{CH}_2 \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \right)$
2342	5- $(-\text{CONH}_2)$	4- $(-\text{NHMe})$
2343	5- $(-\text{CONH}_2)$	4- $(-\text{NHAc})$
2344	5- $(-\text{CONH}_2)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{N} \text{---} \text{S} \text{---} \text{Me} \right)$
2345	5- $(-\text{CONH}_2)$	4- $(-\text{SMe})$
2346	5- $(-\text{CONH}_2)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{Me} \right)$
2347	5- $(-\text{CONH}_2)$	4- $\left( \text{---} \overset{\text{O}}{\parallel} \text{S} \text{---} \text{Me} \right)$



2348	5-(-CONH <sub>2</sub> )	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \text{O} \end{array}\right)$
2349	5-(-CONH <sub>2</sub> )	4- $\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \text{O} \end{array}\right\}$
2350	5-{-CON(Me) <sub>2</sub> }	-H
2351	5-{-CON(Me) <sub>2</sub> }	4-(-F)
2352	4-{-CON(Me) <sub>2</sub> }	4-(-Cl)
2353	5-{-CON(Me) <sub>2</sub> }	4-(-CN)
2354	5-{-CON(Me) <sub>2</sub> }	4-(-NO <sub>2</sub> )
2355	5-{-CON(Me) <sub>2</sub> }	4-(-Me)
2356	5-{-CON(Me) <sub>2</sub> }	4-(-CF <sub>3</sub> )
2357	5-{-CON(Me) <sub>2</sub> }	4-(-Ac)
2358	5-{-CON(Me) <sub>2</sub> }	4-(-CO <sub>2</sub> H)
2359	5-{-CON(Me) <sub>2</sub> }	4-(-CO <sub>2</sub> Me)
2360	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array}\right)$
2361	5-{-CON(Me) <sub>2</sub> }	4-(-CONH <sub>2</sub> )
2362	5-{-CON(Me) <sub>2</sub> }	4-{-CON(Me) <sub>2</sub> }
2363	5-{-CON(Me) <sub>2</sub> }	4-{-C(=NH)NH <sub>2</sub> }
2364	5-{-CON(Me) <sub>2</sub> }	4-(-OMe)
2365	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N} \end{array}\right)$
2366	5-{-CON(Me) <sub>2</sub> }	4-(-NHMe)
2367	5-{-CON(Me) <sub>2</sub> }	4-(-NHAc)
2368	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\ \text{H} \quad \text{O} \end{array}\right)$
2369	5-{-CON(Me) <sub>2</sub> }	4-(-SMe)
2370	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array}\right)$
2371	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \text{O} \end{array}\right)$
2372	5-{-CON(Me) <sub>2</sub> }	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \text{O} \end{array}\right)$

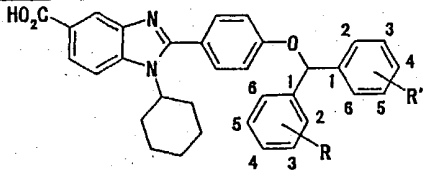
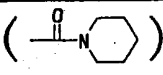
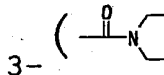
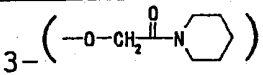
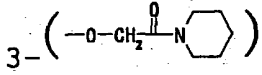
2373	5-{-CON(Me) <sub>2</sub> }	$4-\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$
2374	5-(-OMe)	-H
2375	5-(-OMe)	4-(-F)
2376	5-(-OMe)	4-(-Cl)
2377	5-(-OMe)	4-(-CN)
2378	5-(-OMe)	4-(-NO <sub>2</sub> )
2379	5-(-OMe)	4-(-Me)
2380	5-(-OMe)	4-(-CF <sub>3</sub> )
2381	5-(-OMe)	4-(-Ac)
2382	5-(-OMe)	4-(-CO <sub>2</sub> H)
2383	5-(-OMe)	4-(-CO <sub>2</sub> Me)
2384	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array} \right)$
2385	5-(-OMe)	4-(-CONH <sub>2</sub> )
2386	5-(-OMe)	4-{-CON(Me) <sub>2</sub> }
2387	5-(-OMe)	4-{-C(=NH)NH <sub>2</sub> }
2388	5-(-OMe)	4-(-OMe)
2389	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N} \end{array} \right)$
2390	5-(-OMe)	4-(-NHMe)
2391	5-(-OMe)	4-(-NHAc)
2392	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\ \parallel \\ \text{H} \quad \text{O} \end{array} \right)$
2393	5-(-OMe)	4-(-SMe)
2394	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$
2395	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$
2396	5-(-OMe)	$4-\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$
2397	5-(-OMe)	$4-\left\{ \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \parallel \\ \text{O} \end{array} \right\}$
2398	5-(-NHMe)	4-(-F)

2399	5-(-NHMe)	4-(-Cl)
2400	5-(-NHAc)	4-(-F)
2401	5-(-NHAc)	4-(-Cl)
2402	5-(-NHAc)	4-(-Ac)
2403	5-(-NHAc)	4-(-CONH <sub>2</sub> )
2404	5-(-NHAc)	4-(-CON(Me) <sub>2</sub> )
2405	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-F)
2406	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-Cl)
2407	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-Me)
2408	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-CF <sub>3</sub> )
2409	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-CO <sub>2</sub> H)
2410	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-CO <sub>2</sub> Me)
2411	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--} \end{array}\right)$ 
2412	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4-(-SMe)
2413	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--S--Me} \\   \\ \text{O} \end{array}\right)$
2414	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--N--S--Me} \\   \quad   \\ \text{H} \quad \text{O} \end{array}\right)$	4- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--S--Me} \\   \\ \text{O} \end{array}\right)$
2415	5-(-SMe)	4-(-F)
2416	5-(-SMe)	4-(-Cl)
2417	5-(-SMe)	4-(-Me)
2418	5-(-SMe)	4-(-CF <sub>3</sub> )
2419	5-(-SMe)	4-(-Ac)
2420	5-(-SMe)	4-(-CONH <sub>2</sub> )
2421	5-(-SMe)	4-(-CON(Me) <sub>2</sub> )
2422	5- $\left(\begin{array}{c} \text{O} \\ \parallel \\ \text{--S--Me} \\   \\ \text{O} \end{array}\right)$	4-(-F)

2423	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Cl)
2424	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Me)
2425	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-CF <sub>3</sub> )
2426	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-Ac)
2427	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- (-CONH <sub>2</sub> )
2428	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array} \right)$	4- {-CON (Me) <sub>2</sub> }
2429	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-F)
2430	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-Cl)
2431	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-Me)
2432	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-CF <sub>3</sub> )
2433	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-Ac)
2434	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- (-CONH <sub>2</sub> )
2435	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \parallel \\ \text{O} \end{array} \right)$	4- {-CON (Me) <sub>2</sub> }
2436	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- (-F)
2437	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- (-Cl)
2438	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- (-Me)
2439	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- (-CF <sub>3</sub> )
2440	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- (-CONH <sub>2</sub> )
2441	5- $\left( \begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array} \right)$	4- {-CON (Me) <sub>2</sub> }

2442	$5-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \mid \\ \text{O} \end{array}\right)$	$4-(-\text{SMe})$
2443	$5-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \mid \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \mid \\ \text{O} \end{array}\right)$
2444	$5-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\ \mid \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \mid \\ \text{O} \end{array}\right)$
2445	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{F})$
2446	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{Cl})$
2447	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{Me})$
2448	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{CF}_3)$
2449	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{CONH}_2)$
2450	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{CON}(\text{Me})_2)$
2451	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-(-\text{SMe})$
2452	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \mid \\ \text{O} \end{array}\right)$
2453	$5-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\ \mid \\ \text{O} \end{array}\right\}$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\ \mid \\ \text{O} \end{array}\right)$

Table 215

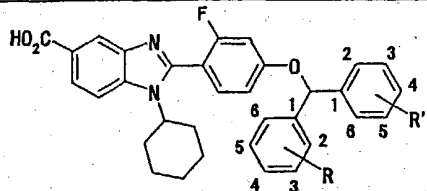
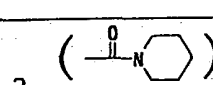
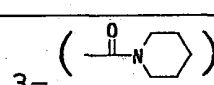
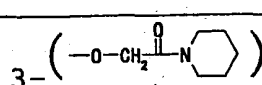
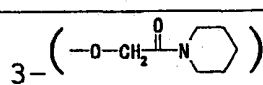
		
Ex. N o.	R	R'
2454	2-(-F)	2-(-F)
2455	2-(-F)	3-(-F)
2456	2-(-F)	4-(-F)
2457	3-(-Cl)	3-(-Cl)
2458	3,5-di-(-Cl)	3,5-di-(-Cl)
2459	3-(-CN)	3-(-CN)
2460	3-(-NO <sub>2</sub> )	3-(-NO <sub>2</sub> )
2461	3-(-Me)	3-(-Me)
2462	3-(-CF <sub>3</sub> )	3-(-CF <sub>3</sub> )
2463	3-(-Ac)	3-(-Ac)
2464	3-(-CO <sub>2</sub> H)	3-(-CO <sub>2</sub> H)
2465	3-(-CO <sub>2</sub> Me)	3-(-CO <sub>2</sub> Me)
2466	3- 	3- 
2467	3-(-CONH <sub>2</sub> )	3-(-CONH <sub>2</sub> )
2468	3-(-CONH <sub>2</sub> )	3-(-F)
2469	3-(-CONH <sub>2</sub> )	3-(-Cl)
2470	3-(-CON(Me) <sub>2</sub> )	3-(-CON(Me) <sub>2</sub> )
2471	3-(-CON(Me) <sub>2</sub> )	3-(-F)
2472	3-(-CON(Me) <sub>2</sub> )	3-(-Cl)
2473	3-(-C(=NH)NH <sub>2</sub> )	3-(-C(=NH)NH <sub>2</sub> )
2474	3-(-OMe)	3-(-OMe)
2475	3- 	3- 
2476	3-(-NHMe)	3-(-NHMe)

2477	3-(-NHAc)	3-(-NHAc)
2478	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \end{array}\right)$
2479	3-(-SMe)	3-(-SMe)
2480	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array}\right)$
2481	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$
2482	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$
2483	$3-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$	$3-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$
2484	3-(-F)	4-(-F)
2485	3-(-Cl)	4-(-Cl)
2486	4-(-CN)	4-(-CN)
2487	4-(-NO <sub>2</sub> )	4-(-NO <sub>2</sub> )
2488	3-(-Me)	4-(-Me)
2489	4-(-Me)	2,6-di-(-Me)
2490	4-(-CF <sub>3</sub> )	4-(-CF <sub>3</sub> )
2491	4-(-Ac)	4-(-Ac)
2492	4-(-CO <sub>2</sub> H)	4-(-CO <sub>2</sub> H)
2493	4-(-CO <sub>2</sub> Me)	4-(-CO <sub>2</sub> Me)
2494	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array} \text{C}_6\text{H}_{11}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N} \end{array} \text{C}_6\text{H}_{11}\right)$
2495	4-(-CONH <sub>2</sub> )	4-(-CONH <sub>2</sub> )
2496	4-(-CONH <sub>2</sub> )	4-(-F)
2497	4-(-CONH <sub>2</sub> )	2,3,4,5,6-penta-(-F)
2498	4-(-CONH <sub>2</sub> )	4-(-Cl)
2499	4-(-CON(Me) <sub>2</sub> )	4-(-CON(Me) <sub>2</sub> )
2500	4-(-CON(Me) <sub>2</sub> )	4-(-F)
2501	4-(-CON(Me) <sub>2</sub> )	4-(-Cl)
2502	4-(-CON(Me) <sub>2</sub> )	3,5-di-(-Cl)
2503	4-(-C(=NH)NH <sub>2</sub> )	4-(-C(=NH)NH <sub>2</sub> )

2504	4-(-OMe)	4-(-OMe)
2505	4-(-OMe)	3,4,5-tri-(-OMe)
2506	$4-\left(-O-CH_2-\overset{\overset{O}{\parallel}}{C}-N\text{C}_6\text{H}_{11}\right)$	$4-\left(-O-CH_2-\overset{\overset{O}{\parallel}}{C}-N\text{C}_6\text{H}_{11}\right)$
2507	4-(-NHMe)	4-(-NHMe)
2508	4-(-NHAc)	4-(-NHAc)
2509	$4-\left(-N\overset{\overset{O}{\parallel}}{\underset{\underset{H}{\mid}}{S}}-Me\right)$	$4-\left(-N\overset{\overset{O}{\parallel}}{\underset{\underset{H}{\mid}}{S}}-Me\right)$
2510	4-(-SMe)	4-(-SMe)
2511	$4-\left(-\overset{\overset{O}{\parallel}}{S}-Me\right)$	$4-\left(-\overset{\overset{O}{\parallel}}{S}-Me\right)$
2512	$4-\left(-\overset{\overset{O}{\parallel}}{S}-Me\right)$	$4-\left(-\overset{\overset{O}{\parallel}}{S}-Me\right)$
2513	$4-\left(-\overset{\overset{O}{\parallel}}{S}-NH_2\right)$	$4-\left(-\overset{\overset{O}{\parallel}}{S}-NH_2\right)$
2514	$4-\left\{-\overset{\overset{O}{\parallel}}{S}-N(Me)_2\right\}$	$4-\left\{-\overset{\overset{O}{\parallel}}{S}-N(Me)_2\right\}$



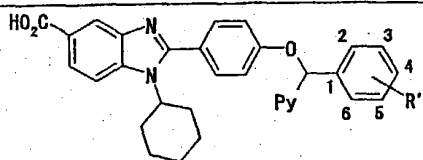
Table 216

		
Ex. N o.	R	R'
2515	-H	-H
2516	2-(-F)	3-(-F)
2517	3-(-Cl)	3-(-Cl)
2518	3-(-CN)	3-(-CN)
2519	3-(-NO <sub>2</sub> )	3-(-NO <sub>2</sub> )
2520	3-(-Me)	3-(-Me)
2521	3-(-CF <sub>3</sub> )	3-(-CF <sub>3</sub> )
2522	3-(-Ac)	3-(-Ac)
2523	3-(-CO <sub>2</sub> H)	3-(-CO <sub>2</sub> H)
2524	3-(-CO <sub>2</sub> Me)	3-(-CO <sub>2</sub> Me)
2525	3- 	3- 
2526	3-(-CONH <sub>2</sub> )	3-(-CONH <sub>2</sub> )
2527	3-(-CONH <sub>2</sub> )	3-(-F)
2528	3-(-CONH <sub>2</sub> )	3-(-Cl)
2529	3-{-CON(Me) <sub>2</sub> }	3-{-CON(Me) <sub>2</sub> }
2530	3-{-CON(Me) <sub>2</sub> }	3-(-F)
2531	3-{-CON(Me) <sub>2</sub> }	3-(-Cl)
2532	3-{-C(=NH)NH <sub>2</sub> }	3-{-C(=NH)NH <sub>2</sub> }
2533	3-(-OMe)	3-(-OMe)
2534	3- 	3- 
2535	3-(-NHMe)	3-(-NHMe)
2536	3-(-NHAc)	3-(-NHAc)

2537	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$
2538	$3-(-\text{SMe})$	$3-(-\text{SMe})$
2539	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \end{array}\right)$
2540	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$
2541	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$	$3-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$
2542	$3-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$	$3-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$
2543	$3-(-\text{F})$	$4-(-\text{F})$
2544	$4-(-\text{Cl})$	$4-(-\text{Cl})$
2545	$4-(-\text{CN})$	$4-(-\text{CN})$
2546	$4-(-\text{NO}_2)$	$4-(-\text{NO}_2)$
2547	$4-(-\text{Me})$	$4-(-\text{Me})$
2548	$4-(-\text{CF}_3)$	$4-(-\text{CF}_3)$
2549	$4-(-\text{Ac})$	$4-(-\text{Ac})$
2550	$3-(-\text{CO}_2\text{H})$	$4-(-\text{CO}_2\text{H})$
2551	$4-(-\text{CO}_2\text{Me})$	$4-(-\text{CO}_2\text{Me})$
2552	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{C}_6\text{H}_{11} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{C}_6\text{H}_{11} \end{array}\right)$
2553	$4-(-\text{CONH}_2)$	$4-(-\text{CONH}_2)$
2554	$4-(-\text{CONH}_2)$	$4-(-\text{F})$
2555	$4-(-\text{CONH}_2)$	$4-(-\text{Cl})$
2556	$3-\{-\text{CON}(\text{Me})_2\}$	$4-\{-\text{CON}(\text{Me})_2\}$
2557	$3-\{-\text{CON}(\text{Me})_2\}$	$4-(-\text{F})$
2558	$4-\{-\text{CON}(\text{Me})_2\}$	$4-(-\text{Cl})$
2559	$4-\{-\text{C}(=\text{NH})\text{NH}_2\}$	$4-\{-\text{C}(=\text{NH})\text{NH}_2\}$
2560	$4-(-\text{OMe})$	$4-(-\text{OMe})$
2561	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N}-\text{C}_6\text{H}_{11} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{CH}_2-\text{N}-\text{C}_6\text{H}_{11} \end{array}\right)$
2562	$4-(-\text{NHMe})$	$4-(-\text{NHMe})$
2563	$4-(-\text{NHAc})$	$4-(-\text{NHAc})$

2564	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{N}-\text{S}-\text{Me} \\   \\ \text{H} \\ \text{O} \end{array}\right)$
2565	$4-(-\text{SMe})$	$4-(-\text{SMe})$
2566	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$
2567	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{Me} \\   \\ \text{O} \end{array}\right)$
2568	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$	$4-\left(\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{NH}_2 \\   \\ \text{O} \end{array}\right)$
2569	$4-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$	$4-\left\{\begin{array}{c} \text{O} \\ \parallel \\ -\text{S}-\text{N}(\text{Me})_2 \\   \\ \text{O} \end{array}\right\}$

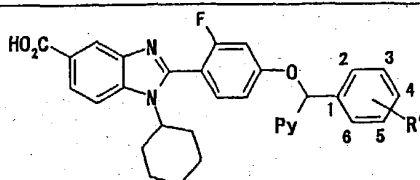
Table 217



Py : pyridyl group

Ex. N o.	Py	R'
2570	3-Py	-H
2571	3-Py	3-(-F)
2572	3-Py	3-(-Cl)
2573	3-Py	3-(-Me)
2574	3-Py	3-(-CF <sub>3</sub> )
2575	3-Py	3-(-Ac)
2576	3-Py	3-(-CO <sub>2</sub> H)
2577	3-Py	3-(-CO <sub>2</sub> Me)
2578	3-Py	3-
2579	3-Py	3-(-CONH <sub>2</sub> )
2580	3-Py	3-{-CON(Me) <sub>2</sub> }
2581	3-Py	4-(-F)
2582	3-Py	4-(-Cl)
2583	3-Py	4-(-Me)
2584	3-Py	4-(-CF <sub>3</sub> )
2585	3-Py	4-(-Ac)
2586	2-Py	4-(-CO <sub>2</sub> H)
2587	3-Py	4-(-CO <sub>2</sub> Me)
2588	3-Py	4-
2589	4-Py	4-(-CONH <sub>2</sub> )
2590	3-Py	4-{-CON(Me) <sub>2</sub> }

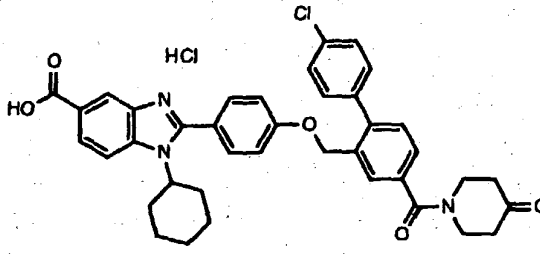
Table 218

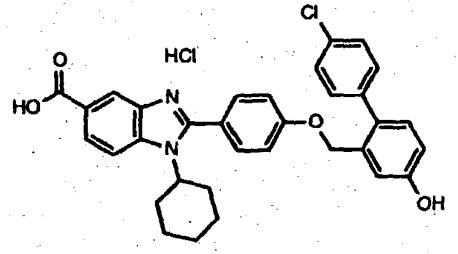


Py : pyridyl group

Ex. No.	Py	R'
2591	3-Py	-H
2592	3-Py	3-(-F)
2593	3-Py	3-(-Cl)
2594	3-Py	3-(-Me)
2595	3-Py	3-(-CF <sub>3</sub> )
2596	3-Py	3-(-Ac)
2597	3-Py	3-(-CO <sub>2</sub> H)
2598	3-Py	3-(-CO <sub>2</sub> Me)
2599	3-Py	3-
2600	3-Py	3-(-CONH <sub>2</sub> )
2601	3-Py	3-{-CON(Me) <sub>2</sub> }
2602	3-Py	4-(-F)
2603	3-Py	4-(-Cl)
2604	3-Py	4-(-Me)
2605	3-Py	4-(-CF <sub>3</sub> )
2606	3-Py	4-(-Ac)
2607	3-Py	4-(-CO <sub>2</sub> H)
2608	3-Py	4-(-CO <sub>2</sub> Me)
2609	3-Py	4-
2610	3-Py	4-(-CONH <sub>2</sub> )
2611	3-Py	4-{-CON(Me) <sub>2</sub> }

Table 219

Example No.	328	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.23 (1H, d, J=9.0 Hz), 8.02 (1H, d, J=8.4Hz), 7.80 (1H, s), 7.71 (2H, d, J=8.4Hz), 7.61 (1H, d, J=9.3Hz), 7.55-7.45 (3H, m), 7.46 (2H, d, J=8.1Hz), 7.22 (2H, d, J=8.7Hz), 5.16 (2H, s), 4.34 (1H, m), 4.20-3.40 (4H, m), 2.60-2.15 (6H, m), 2.10-1.90 (2H, m), 1.85-1.70 (2H, m), 1.65-1.55 (1H, m), 1.50-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	662 (M+1)	

Example No.	329	1H NMR (δ) ppm
		400MHz, DMSO-d6 9.80 (1H, brs), 8.32 (1H, s), 8.30 (1H, d, J=8.8Hz), 8.06 (1H, d, J=8.8Hz), 7.74 (2H, d, J=8.6Hz), 7.48-7.37 (4H, m), 7.22 (1H, d, J=8.6Hz), 7.17 (1H, d, J=8.2Hz), 7.05 (1H, d, J=2.3Hz), 6.88 (1H, dd, J=8.3, 2.5Hz), 5.04 (2H, s), 4.37 (1H, m), 2.37-2.22 (2H, m), 2.11-1.98 (2H, m), 1.93-1.81 (2H, m), 1.70-1.58 (1H, m), 1.56-1.22 (3H, m)
Purity	> 90% (NMR)	
MS	553 (M+1)	

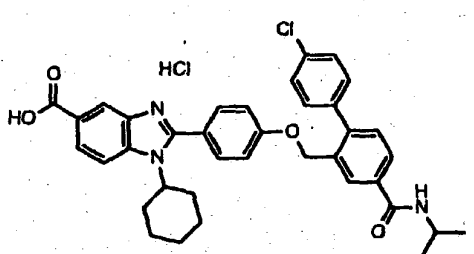
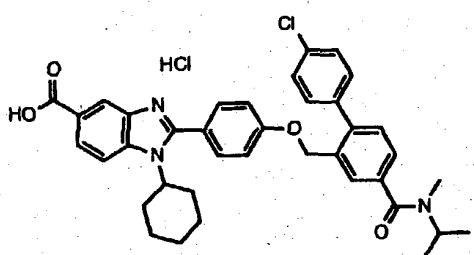
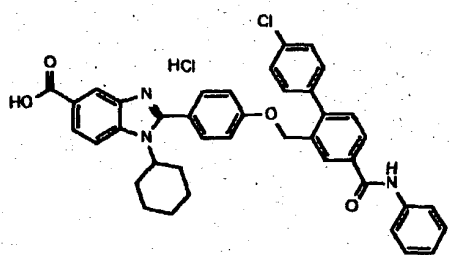
Example No.	330	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.38 (1H, d, J=7.5Hz), 8.32 (1H, s), 8.29 (1H, d, J=9.0Hz), 8.16 (1H, s), 8.05 (1H, d, J=9.0Hz), 7.96 (1H, d, J=7.5Hz), 7.75 (2H, d, J=8.4Hz), 7.53-7.43 (5H, m), 7.25 (2H, d, J=8.4Hz), 5.13 (2H, s), 4.36 (1H, m), 4.12 (1H, sept, J=6.9Hz), 2.40-2.15 (2H, m), 2.10-1.95 (2H, m), 1.90-1.75 (2H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m), 1.18 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	622 (M+1)	

Table 220

Example No.	331	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.27 (1H, d, J=8.7Hz), 8.05 (1H, d, J=8.7Hz), 7.75-7.41 (9H, m), 7.23 (2H, d, J=8.7Hz), 4.36 (1H, m), 4.00-3.90 (1H, m), 2.84 (3H, brs), 2.40-2.15 (2H, m), 2.10-2.00 (2H, m), 1.95-1.75 (2H, m), 1.70-1.55 (1H, m), 1.50-1.00 (7H, m)
Purity	> 90% (NMR)	
MS	636 (M+1)	

Example No.	332	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.42 (1H, s), 8.29 (1H, s), 8.27 (1H, s), 8.10 (1H, d, J=7.9Hz), 8.03 (1H, d, J=8.6Hz), 7.82 (2H, d, J=7.5Hz), 7.73 (2H, d, J=8.7Hz), 7.56-7.52 (5H, m), 7.38 (2H, t, J=7.9Hz), 7.26 (2H, d, J=8.7Hz), 7.13 (1H, t, J=7.5Hz), 5.20 (2H, s), 4.35 (1H, br t, J=11.7Hz), 2.37-2.19 (2H, m), 2.07-1.96 (2H, m), 1.92-1.79 (2H, m), 1.69-1.58 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	656 (M+1)	

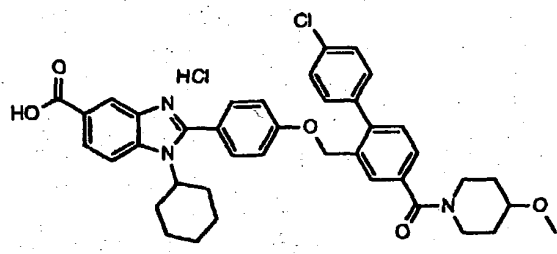
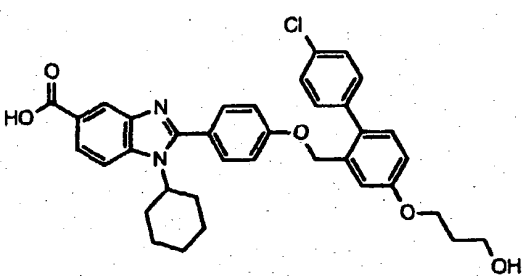
Example No.	333	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.24 and 8.03 (2H, ABq, J=8.8Hz), 7.71 and 7.22 (4H, A'B'q, J=8.8Hz), 7.69 (1H, s), 7.52 (4H, s), 7.50 and 7.43 (2H, A''B''q, J=7.7Hz), 5.15 (2H, s), 4.35 (1H, br t, J=12.1Hz), 4.05-3.15 (5H, br m), 3.27 (3H, s), 2.39-2.20 (2H, m), 2.07-1.75 (6H, m), 1.70-1.58 (1H, m), 1.55-1.20 (5H, m)
Purity	> 90% (NMR)	
MS	678 (M+1)	

Table 221

Example No.	334	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.22 (1H, d, J=1.5Hz), 8.01 (1H, d, J=9.0Hz), 7.89 (1H, dd, J=8.6, 1.5Hz), 7.61 (2H, d, J=8.6Hz), 7.50-7.39 (4H, m), 7.27 (1H, d, J=8.6Hz), 7.22 (1H, d, J=2.6Hz), 7.13 (2H, d, J=8.6Hz), 7.04 (1H, dd, J=8.2, 2.6Hz), 5.04 (2H, s), 4.28 (1H, m), 4.11 (2H, t, J=6.3Hz), 3.57 (2H, t, J=6.3Hz), 2.38-2.17 (2H, m), 2.00-1.79 (6H, m), 1.70-1.59 (1H, m), 1.52-1.16 (3H, m)
Purity	> 90% (NMR)	
MS	611 (M+1)	

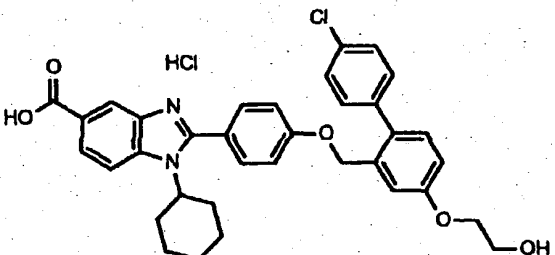
Example No.	335	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, d, J=1.5Hz), 8.27 (1H, d, J=9.0Hz), 8.04 (1H, dd, J=8.6, 1.5Hz), 7.72 (2H, d, J=9.0Hz), 7.60-7.40 (4H, m), 7.32-7.19 (4H, m), 7.06 (1H, dd, J=8.6, 3.0Hz), 5.08 (2H, s), 4.36 (1H, m), 4.06 (2H, t, J=4.8Hz), 3.74 (2H, t, J=4.8Hz), 2.38-2.19 (2H, m), 2.13-1.97 (2H, m), 1.94-1.78 (2H, m), 1.72-1.59 (1H, m), 1.52-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	597 (M+1)	



Table 222

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
340	0.017	360	0.014
341	0.025	361	0.028
342	0.015	362	0.020
343	0.017	363	0.11
344	0.016	364	0.12
345	0.012	365	0.020
346	0.025	366	0.024
347	0.022	367	0.011
348	0.013	368	0.024
349	0.021	369	0.022
350	0.020	370	0.017
351	0.019	371	0.015
352	0.013	372	0.033
353	0.023	373	0.013
354	0.013	374	0.013
355	0.015	375	0.012
356	0.016	376	0.014
357	0.019	377	0.012
358	0.017	378	0.018
359	0.015	379	0.021

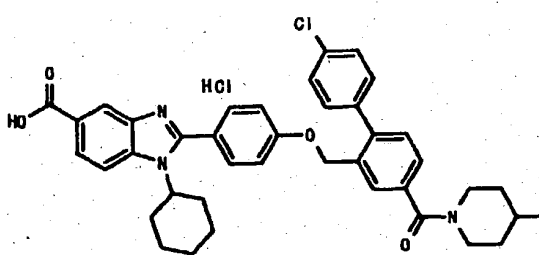
Table 223

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [ $\mu$ M]
380	0.023	409	0.020
381	0.011	410	0.018
382	0.015	411	0.015
383	0.013	412	0.019
384	0.016	413	0.026
385	0.019	414	0.024
386	0.018	415	0.019
387	0.025	416	0.024
388	0.020	417	0.029
389	0.012	418	0.016
390	0.014	419	0.021
391	0.017	420	0.015
392	0.014	421	0.017
393	0.011	422	0.017
394	0.019	423	0.017
395	0.016	424	0.020
396	0.025	425	0.026
397	0.037	426	0.053
398	0.077	427	0.020
399	0.032	428	0.026

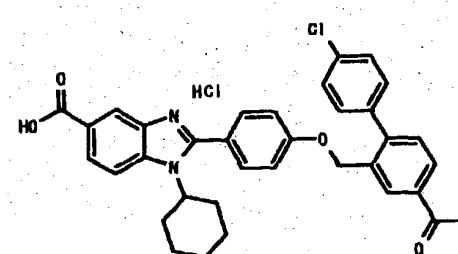
Table 224

Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]	Ex. No.	HCV polymerase inhibitory activity IC <sub>50</sub> [μM]
429	0.017	442	0.024
430	0.017	443	0.030
431	0.015	445	0.33
432	0.022	446	0.016
433	0.014	502	0.024
434	0.011	503	0.196
435	0.012	601	0.32
436	0.026	701	0.052
440	0.070		

Table 225

Example No.	341	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.29 (1H, d, J=1.5Hz), 8.25 (1H, d, J=8.7Hz), 8.03 (1H, dd, J=8.7Hz), 7.72 and 7.22 (4H, Abq, J=8.8Hz), 7.67 (1H, d, J=1.5Hz), 7.52 (4H, s), 7.49 (1H, dd, J=7.9, 1.5Hz), 7.43 (1H, d, J=7.9Hz), 4.46 (1H, brs), 4.35 (1H, brt, J=12.4Hz), 3.62 (1H, brs), 3.06 (1H, brs), 2.79 (1H, brs), 2.38-2.20 (2H, brm), 2.08-1.81 (4H, brm), 1.77-1.52 (4H, brm), 1.46-1.20 (3H, brm), 1.19-1.00 (2H, brm), 0.94 and 0.92 (total 13H, each s)
Purity	> 90% (NMR)	
MS	662 (M+1)	

Example No.	342	1H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.28 (1H, d, J=1.5Hz), 8.26 (1H, d, J=1.8Hz), 8.19 (1H, d, J=8.8Hz), 8.07 (1H, dd, J=7.7, 1.8Hz), 8.00 (1H, dd, J=8.8, 1.5Hz), 7.70 and 7.22 (4H, Abq, J=8.8Hz), 7.56-7.50 (1H, m), 7.56 (4H, s), 5.17 (2H, s), 4.33 (1H, brt, J=12.5Hz), 2.05 (3H, s), 2.37-2.20 (2H, brm), 2.06-1.80 (4H, brm), 1.70-1.60 (1H, brm), 1.50-1.20 (3H, brm)
Purity	> 90% (NMR)	
MS	679 (M+1)	

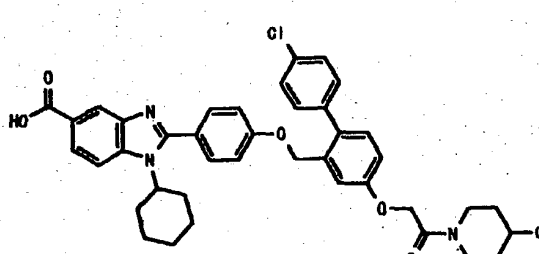
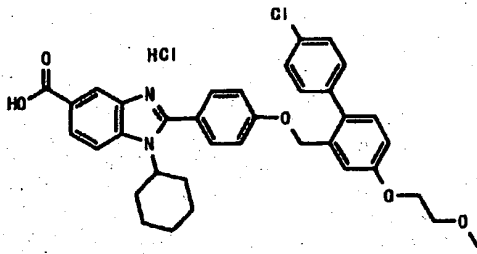
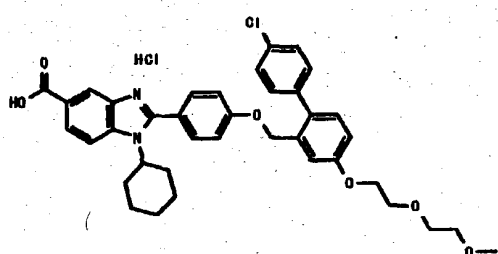
Example No.	343	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.20 (1H, d, J=1.5Hz), 7.93 (1H, d, J=8.6Hz), 7.84 (1H, dd, J=8.3Hz, 1.5Hz), 7.57 (2H, d, J=8.6Hz), 7.50-7.40 (4H, m), 7.27 (1H, d, J=8.2Hz), 7.22 (1H, d, J=2.6Hz), 7.10 (2H, d, J=8.6Hz), 7.01 (1H, dd, J=8.6Hz, 2.6Hz), 5.02 (2H, s), 4.89 (2H, s), 4.78 (1H, d, J=4.1Hz), 4.38-4.18 (1H, m), 3.96-3.81 (1H, m), 3.78-3.62 (2H, m), 3.27-2.99 (2H, m), 2.35-1.15 (4H, m)
Purity	> 90% (NMR)	
MS	694 (M+1)	

Table 226

Example No.	344	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 8.23 (1H, d, J=8.7Hz), 8.02 (1H, d, J=8.4Hz), 7.71 (2H, d, J=8.7Hz), 7.55-7.15 (8H, m), 7.07 (1H, dd, J=8.4Hz, 3.0Hz), 5.07 (2H, s), 4.35 (1H, m), 4.17 (2H, t, J=4.5Hz), 3.69 (2H, t, J=4.5Hz), 3.32 (3H, s), 2.40-2.15 (2H, m), 2.10-1.80 (4H, m), 1.75-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	611 (M+1)	

Example No.	345	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.29 (1H, d, J=1.5Hz), 8.22 (1H, d, J=8.7Hz), 8.01 (1H, d, J=8.7Hz), 7.70 (1H, d, J=8.7Hz), 7.50-7.15 (8H, m), 7.07 (1H, dd, J=8.4Hz, 2.4Hz), 5.07 (2H, s), 4.35 (1H, m), 4.17 (2H, t, J=4.2Hz), 3.76 (2H, t, J=4.5Hz), 3.65-3.40 (4H, m), 3.25 (3H, s), 2.40-2.20 (2H, m), 2.10-1.80 (4H, m), 1.75-1.65 (1H, m), 1.65-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	655 (M+1)	

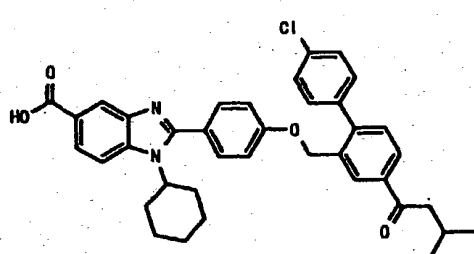
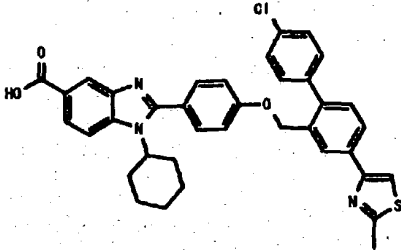
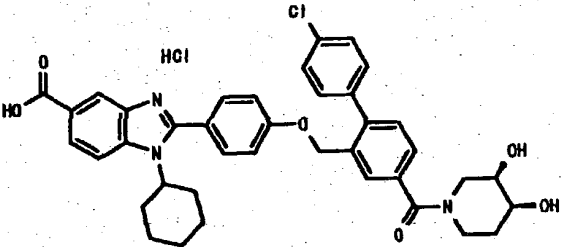
Example No.	346	1H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.26 (1H, d, J=1.9Hz), 8.23 (1H, d, J=1.5Hz), 8.08-8.02 (2H, m), 7.91 (1H, dd, J=8.7, 1.5Hz), 7.63 and 7.16 (4H, Abq, J=8.9Hz), 7.56-7.51 (5H, m), 5.15 (2H, s), 4.29 (1H, brt, J=11.7Hz), 2.96 (2H, d, J=6.9Hz), 2.37-2.12 (3H, m), 2.00-1.79 (4H, brm), 1.71-1.60 (1H, brm), 1.49-1.19 (3H, brm), 0.97 and 0.95 (total 6H, each s)
Purity	> 90% (NMR)	
MS	621 (M+1)	

Table 227

Example No.	347	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.26 (1H, s), 8.22 (1H, s), 8.06 (1H, s), 8.05 (1H, d, J=8.0Hz), 7.94 and 7.85 (2H, ABq, J=8.8Hz), 7.59 and 7.15 (4H, A'B'q, J=8.6Hz), 7.52 (4H, s), 7.44 (1H, d, J=8.0Hz), 5.12 (2H, s), 4.27 (1H, brt, J=11.4Hz), 2.38-2.18 (2H, brm), 1.97-1.77 (4H, brm), 1.70-1.59 (1H, brm), 1.49-1.17 (3H, brm)
Purity	> 90% (NMR)	
MS	634 (M+1)	

Example No.	348	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.29 (1H, d, J=9.0Hz), 8.06 (1H, d, J=8.7Hz), 7.74 (2H, d, J=9.0Hz), 7.72 (1H, brs), 7.60-7.45 (5H, m), 7.42 (1H, d, J=7.8Hz), 7.24 (2H, d, J=8.7Hz), 5.15 (2H, s), 4.37 (1H, m), 4.00-3.10 (6H, m), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.20 (6H, m)
Purity	> 90% (NMR)	
MS	680 (M+1)	

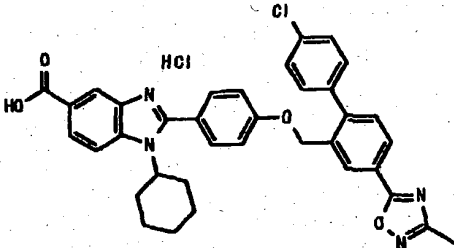
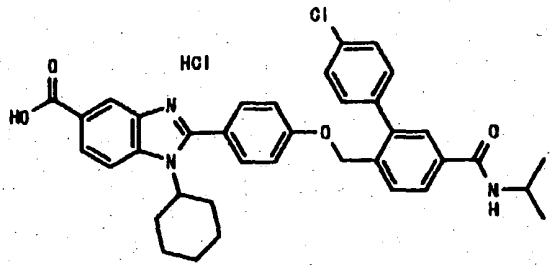
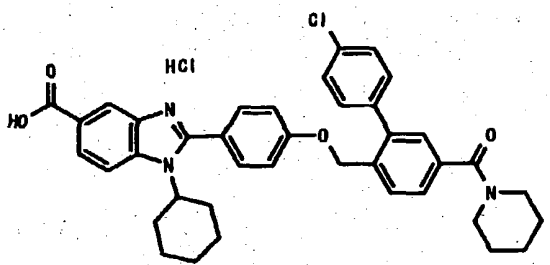
Example No.	349	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.41 (1H, d, J=1.5Hz), 8.33 (1H, d, J=1.5Hz), 8.26 (1H, d, J=8.7Hz), 8.18 (1H, dd, J=2.0Hz, 8.0Hz), 8.04 (1H, dd, J=1.5Hz, 9.0Hz), 7.75 (2H, d, J=8.7Hz), 7.63 (1H, d, J=8.1Hz), 7.62-7.45 (4H, m), 7.26 (2H, d, J=8.7Hz), 5.25 (2H, s), 4.35 (1H, m), 2.45 (3H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	619 (M+1)	

Table 228

Example No.	350	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.36 (1H, d, J=7.7Hz), 8.29 (1H, s), 8.23 (1H, d, J=8.8Hz), 8.02 (1H, d, J=8.6Hz), 7.94 (1H, d, J=7.9Hz), 7.84 (1H, d, J=1.6Hz), 7.80-7.65 (3H, m), 7.53 (4H, s), 5.15 (2H, s), 4.34 (1H, m), 4.12 (1H, m), 2.35-2.20 (2H, m), 2.10-1.60 (5H, m), 1.50-1.20 (3H, m), 1.17 (6H, d, J=6.5Hz)
Purity	> 90% (NMR)	
MS	622 (M+1)	

Example No.	351	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.29 (1H, s), 8.24 (1H, d, J=8.8Hz), 8.02 (1H, d, J=8.6Hz), 7.80-7.65 (3H, m), 7.55-7.45 (5H, m), 7.32 (1H, d, J=1.5Hz), 7.22 (2H, d, J=8.8Hz), 5.13 (2H, s), 4.35 (1H, m), 3.60 (2H, m), 3.33 (2H, m), 2.40-2.15 (2H, m), 2.10-1.15 (14H, m)
Purity	> 90% (NMR)	
MS	648 (M+1)	

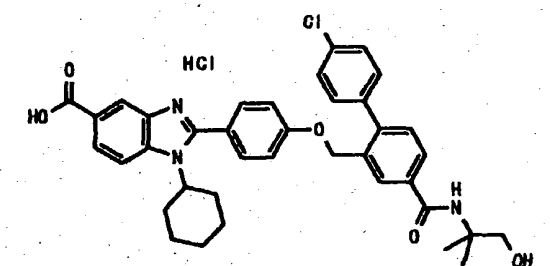
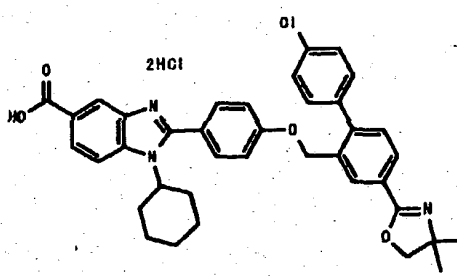
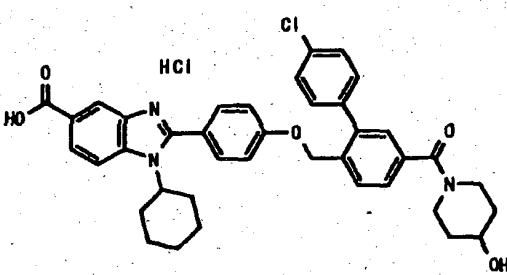
Example No.	352	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.20 (1H, brs), 8.30-8.24 (2H, m), 8.13 (1H, s), 8.04 (1H, d, J=8.7Hz), 7.94 (1H, d, J=8.0Hz), 7.75-7.70 (3H, m), 7.55-7.43 (5H, m), 7.25 (2H, d, J=8.7Hz), 5.13 (2H, s), 4.36 (1H, m), 3.53 (2H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (9H, m)
Purity	> 90% (NMR)	
MS	652 (M+1)	

Table 229

Example No.	353	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.41 (1H, s), 8.33-8.29 (2H, m), 8.16 (1H, d, J=8.2Hz), 8.07 (1H, d, J=8.6Hz), 7.77 (2H, d, J=8.7H z), 7.62 (1H, d, J=8.0Hz), 7.59- 7.51 (4H, m), 7.28 (2H, d, J=8.8H z), 5.21 (2H, s), 4.56 (2H, s), 4. 37 (1H, m), 2.40-2.18 (2H, m), 2. 15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.2 0 (9H, m)
Purity	約90% (NMR)	
MS	634 (M+1)	

Example No.	354	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.25 (1H, d, J=9.0H z), 8.03 (1H, d, J=8.7Hz), 7.76- 7.71 (3H, m), 7.51-7.47 (5H, m), 7.33 (1H, s), 7.23 (2H, d, J=9.0H z), 5.14 (2H, s), 4.36 (1H, m), 4. 02 (1H, m), 3.75 (1H, m), 3.56 (1H , m), 3.22 (2H, m), 2.40-2.18 (2H , m), 2.15-1.95 (2H, m), 1.90-1. 55 (5H, m), 1.50-1.20 (5H, m)
Purity	>90% (NMR)	
MS	664 (M+1)	

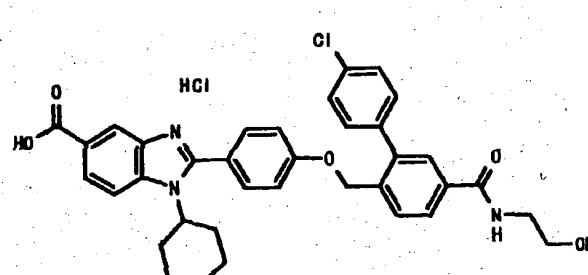
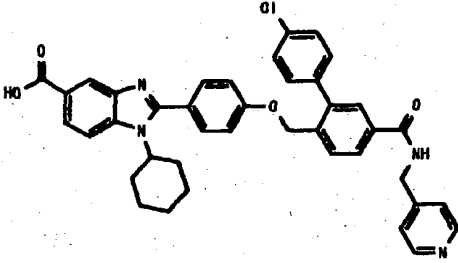
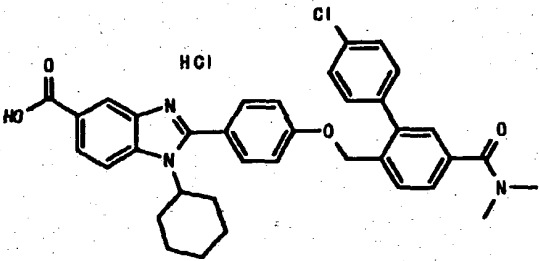
Example No.	355	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.62 (1H, t, J=5.7Hz), 8.32-8.3 0 (2H, m), 8.25 (1H, d, J=8.7Hz), 8.03 (1H, d, J=8.7Hz), 7.96 (1H, d, J=8.1Hz), 7.86 (1H, s), 7.75 ( 1H, d, J=9.0Hz), 7.72 (2H, d, J=9 .0Hz), 7.55-7.50 (4H, m), 7.22 ( 2H, d, J=9.0Hz), 5.17 (2H, s), 4. 35 (1H, m), 3.52 (2H, t, J=6.0Hz) , 3.36 (2H, t, J=6.0Hz), 2.40-2. 18 (2H, m), 2.15-1.95 (2H, m), 1. 90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	>90% (NMR)	
MS	624 (M+1)	



Table 230

Example No.	356	1H NMR (δ) ppm
		300Mz, DMSO-d6 9.30 (1H, t, J=5.9Hz), 8.54 (2H, d, J=5.9Hz), 8.22 (1H, s), 8.02-7.79 (5H, m), 7.59 and 7.12 (4H, ABq, J=8.6Hz), 7.55 (4H, s), 7.37 (2H, d, J=5.9Hz), 5.15 (2H, s), 4.54 (2H, d, J=5.7Hz), 4.26 (1H, brt, J=12.8Hz), 2.36-2.18 (2H, brm), 1.97-1.78 (4H, brm), 1.70-1.60 (1H, brm), 1.47-1.17 (3H, brm)
Purity	> 90% (NMR)	
MS	671 (M+1)	

Example No.	357	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.31 (1H, d, J=1.5Hz), 8.43 (1H, d, J=8.4Hz), 8.03 (1H, dd, J=8.4, 1.5Hz), 7.74 (1H, d, J=8.1Hz), 7.73 and 7.23 (4H, ABq, J=9.0Hz), 7.54-7.51 (5H, m), 7.37 (1H, d, J=1.8Hz), 5.14 (2H, s), 4.36 (1H, brt, J=12.1Hz), 2.98 (6H, brs), 2.37-2.20 (2H, brm), 2.08-1.81 (4H, brm), 1.70-1.60 (1H, brm), 1.50-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	608 (M+1)	

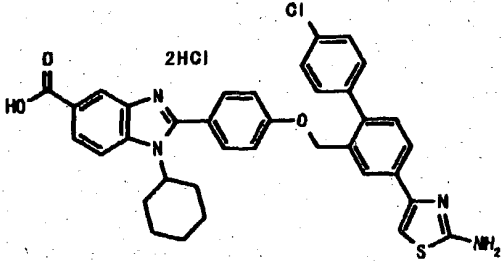
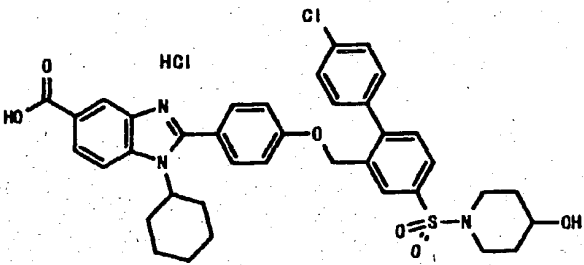
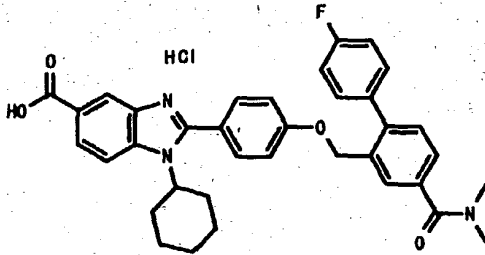
Example No.	358	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.31 (1H, d, J=8.7Hz), 8.14 (1H, s), 8.07 (1H, d, J=8.7Hz), 7.92 (1H, d, J=8.0Hz), 7.76 (2H, d, J=8.7Hz), 7.52-7.40 (5H, m), 7.31-7.26 (3H, m), 5.15 (2H, s), 4.37 (1H, m), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	約90% (NMR)	
MS	635 (M+1)	

Table 231

Example No.	359	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.25 (1H, d, J=8.7Hz), 8.10-7.90 (2H, m), 7.82 (1H, dd, J=7.8Hz, 1.8Hz), 7.72 (2H, d, J=9.0Hz), 7.63 (1H, d, J=8.1Hz), 7.23 (2H, d, J=9.0Hz), 5.25 (2H, s), 4.34 (1H, m), 3.65-3.50 (1H, m), 3.20-3.05 (2H, m), 2.90-2.75 (2H, m), 2.40-2.15 (2H, m), 2.10-1.10 (12H, m)
Purity	> 90% (NMR)	
MS	700 (M+1)	

Example No.	360	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.33 (1H, s), 8.30 (1H, d, J=8.5Hz), 8.06 (1H, d, J=10.1Hz), 8.80-8.65 (3H, m), 8.60-8.45 (3H, m), 7.42 (1H, d, J=7.8Hz), 7.35-7.15 (4H, m), 5.15 (2H, s), 4.36 (1H, m), 3.01, 2.97 (6H, s), 2.40-2.15 (2H, m), 2.10-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	592 (M+1)	

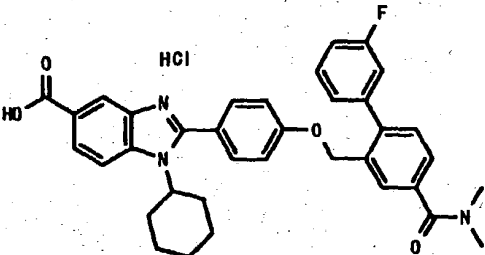
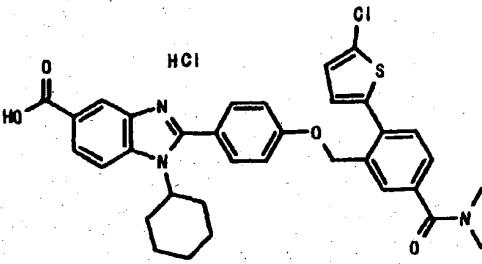
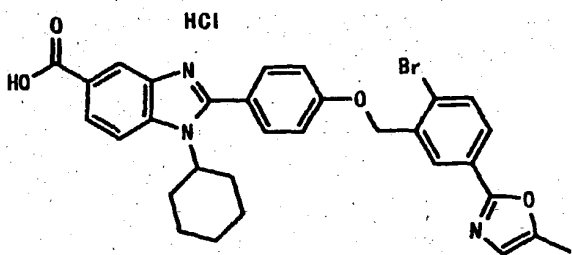
Example No.	361	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.35-8.20 (2H, m), 8.05 (1H, d, J=8.7Hz), 8.80-8.65 (3H, m), 7.60-7.40 (3H, m), 7.40-7.30 (5H, m), 5.17 (2H, s), 4.35 (1H, m), 3.01, 2.97 (6H, s), 2.40-2.15 (2H, m), 2.10-1.80 (4H, m), 1.70-1.20 (4H, m)
Purity	> 90% (NMR)	
MS	592 (M+1)	

Table 232

Example No.	362	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.33 (1H, s), 8.29 (1H, d, J=8.7Hz), 8.06 (1H, d, J=8.7Hz), 7.79 (2H, d, J=9.0Hz), 7.76 (1H, d, J=9.0Hz), 7.60 (1H, d, J=8.1Hz), 7.53 (1H, dd, J=1.7Hz, 8.0Hz), 7.35 (2H, d, J=8.7Hz), 6.85-6.80 (2H, m), 5.29 (2H, s), 4.38 (1H, m), 3.01, 2.96 (6H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	614 (M+1)	

Example No.	363	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.28 (1H, d, J=1.3Hz), 8.20-8.10 (2H, m), 8.98 (1H, d, J=8.6Hz), 7.90-7.80 (2H, m), 7.75 (2H, d, J=8.7Hz), 7.36 (2H, d, J=8.7Hz), 7.04 (1H, d, J=1.3Hz), 5.35 (2H, s), 4.36 (1H, m), 2.39 (3H, s), 2.35-2.15 (2H, m), 2.05-1.75 (4H, m), 1.70-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	586 (M+1)	

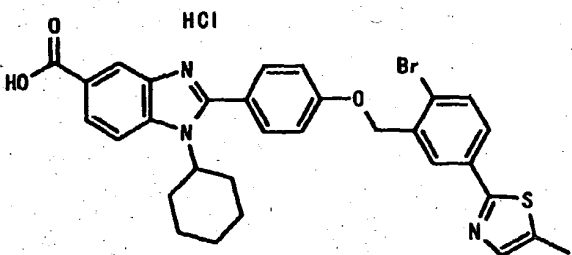
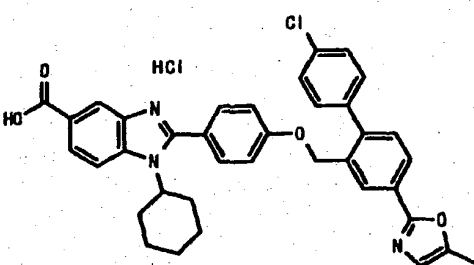
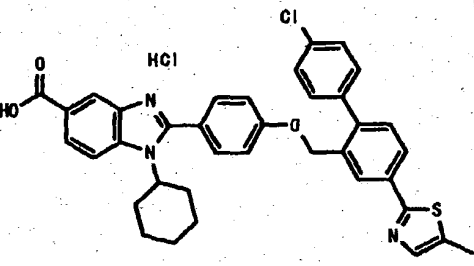
Example No.	364	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31 (1H, s), 8.26 (1H, d, J=8.7Hz), 8.13 (1H, s), 8.04 (1H, d, J=9.0Hz), 7.90-7.70 (4H, m), 7.65 (1H, s), 7.39 (2H, d, J=9.0Hz), 5.37 (2H, s), 4.38 (1H, m), 2.40-2.20 (2H, m), 2.15-2.00 (2H, m), 1.95-1.80 (2H, m), 1.75-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	604 (M+1)	

Table 233

Example No.	365	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.28 (1H, s), 8.23 (1H, s), 8.17 (1H, d, J=8.7Hz), 8.00 (2H, t, J=6.9Hz), 7.69 (2H, d, J=8.4Hz), 7.60-7.45 (5H, m), 7.21 (2H, d, J=8.4Hz), 7.05 (1H, s), 5.19 (2H, s), 4.33 (1H, m), 2.41 (3H, s), 2.40-2.20 (2H, m), 2.10-1.80 (4H, m), 1.70-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	618 (M+1)	

Example No.	366	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.26 (1H, s), 8.17 (1H, s), 8.11 (1H, d, J=8.7Hz), 7.95 (2H, d, J=9.6Hz), 7.70-7.40 (8H, m), 7.19 (2H, d, J=8.4Hz), 5.18 (2H, s), 4.30 (1H, m), 2.51 (3H, s), 2.40-2.15 (2H, m), 2.05-1.80 (4H, m), 1.75-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	634 (M+1)	

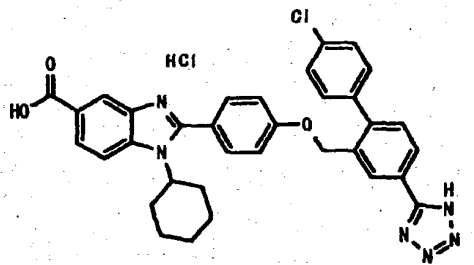
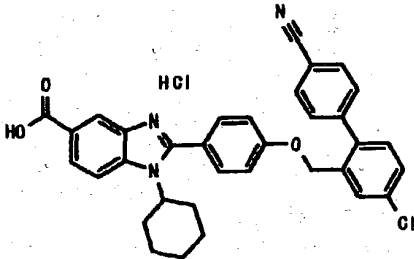
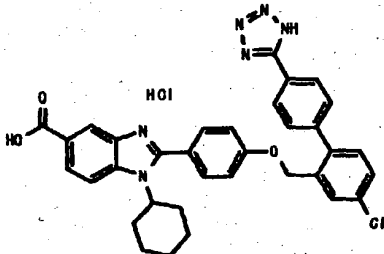
Example No.	367	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.42 (1H, d, J=1.9Hz), 8.30 (1H, J=1.5Hz), 8.27 (1H, d, J=8.7Hz), 8.18 (1H, dd, J=7.9, 1.9Hz), 8.04 (1H, dd, J=8.7, 1.5Hz), 7.75 and 7.29 (4H, ABq, J=8.9Hz), 7.63 (1H, d, J=7.9Hz), 5.23 (2H, s), 4.36 (1H, brt, J=12.3Hz), 2.37-2.20 (2H, brm), 2.08-1.80 (4H, brm), 1.71-1.60 (1H, brm), 1.51-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	605 (M+1)	

Table 234

Example No.	368	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.30 (1H, d, J=1.5Hz), 8.25 (1H, d, J=8.6Hz), 8.04 (1H, dd, J=8.6, 1.5Hz), 7.93 and 7.67 (4H, ABq, J=8.1Hz), 7.80 (1H, d, J=2.2Hz), 7.72 and 7.21 (4H, A'B'q, J=8.6Hz), 7.60 (1H, dd, J=8.1, 2.2Hz), 7.44 (1H, d, J=8.1Hz), 5.13 (2H, s), 4.34 (1H, brt, J=11.7Hz), 2.37-2.19 (2H, brm), 2.09-1.80 (4H, brm), 1.72-1.60 (1H, brm), 1.50-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	562 (M+1)	

Example No.	369	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.30 (1H, d, J=1.5Hz), 8.25 (1H, d, J=8.6Hz), 8.16 and 7.72 (4H, ABq, J=8.4Hz), 8.13 (1H, dd, J=8.6, 1.5Hz), 7.80 (1H, d, J=2.2Hz), 7.70 and 7.24 (4H, A'B'q, J=8.8Hz), 7.61 (1H, dd, J=8.1, 2.2Hz), 7.48 (1H, d, J=8.1Hz), 5.17 (2H, s), 4.33 (1H, brt, J=12.1Hz), 2.36-2.18 (2H, brm), 2.08-1.77 (4H, brm), 1.69-1.57 (1H, brm), 1.49-1.17 (3H, brm)
Purity	> 90% (NMR)	
MS	605 (M+1)	

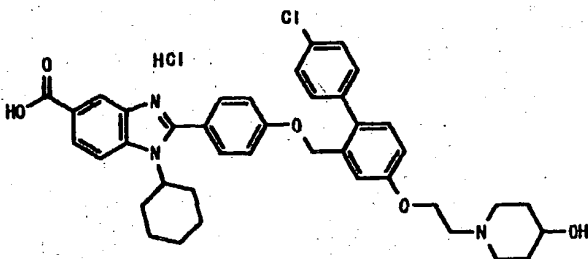
Example No.	370	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.94 (1H, brs), 8.33 (1H, s), 8.27 (1H, d, J=8.7Hz), 8.04 (1H, d, J=8.7Hz), 7.74 (2H, d, J=8.4Hz), 7.56-7.29 (6H, m), 7.23 (2H, d, J=8.7Hz), 7.13 (1H, d, J=8.7Hz), 5.08 (2H, s), 4.51 (2H, brs), 4.36 (1H, m), 3.94 (1H, brs), 3.75-3.00 (6H, m), 3.20-1.20 (14H, m)
Purity	> 90% (NMR)	
MS	680 (M+1)	

Table 235

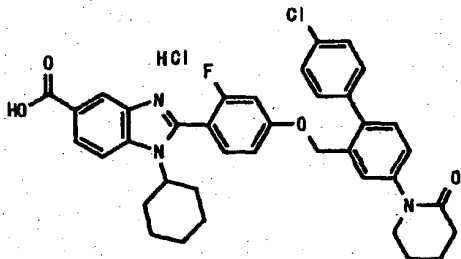
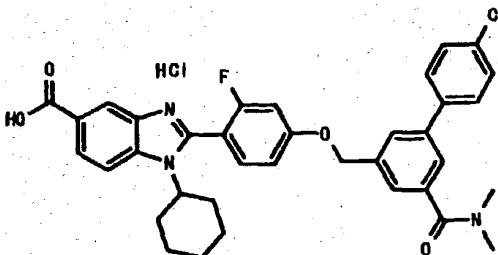
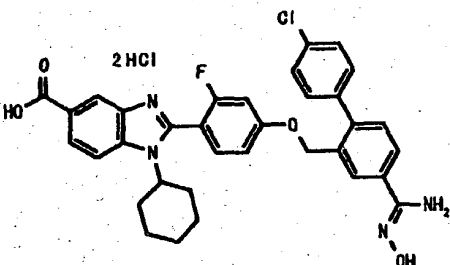
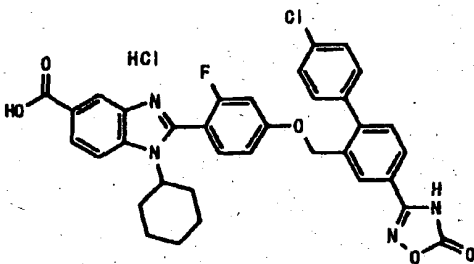
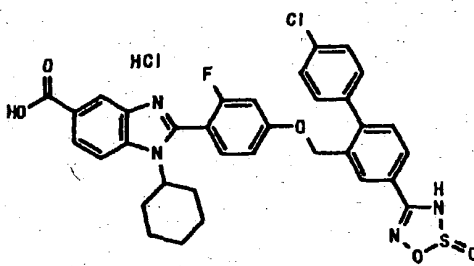
Example No.	371	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31 (1H, d, J=1.5Hz), 8.17 (1H, d, J=9.0Hz), 7.99 (1H, dd, J=8.7 Hz, 1.4Hz), 7.70-7.55 (2H, m), 7.50-7.30 (6H, m), 7.19 (1H, dd, J=12.0Hz, 2.2Hz), 7.06 (1H, dd, J=8.6Hz, 2.2Hz), 5.08 (2H, 4.10 (1H, m), 3.68 (2H, brt, J=5.2), 2.50 (2H, brt, J=1.8Hz), 2.30-2.10 (2H, m), 2.00-1.75 (8H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	652 (M+1)	
Example No.	372	1H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.29 (1H, d, J=1.5Hz), 8.11 (1H, d, J=8.6Hz), 7.96 (1H, dd, J=8.6, 1.5Hz), 7.89 (1H, s), 7.78 and 7.56 (4H, ABq, J=8.4Hz), 7.69 (1H, s), 7.66 (1H, t, J=8.8Hz), 7.31 (1H, dd, J=12.1, 2.2Hz), 7.18 (1H, dd, J=8.8, 2.2Hz), 5.37 (2H, s), 4.08 (1H, brt, J=11.0Hz), 3.02 (3H, s), 2.96 (3H, s), 2.31-2.14 (2H, brm), 1.95-1.77 (4H, brm), 1.69-1.59 (31H, brm), 1.46-1.18 (3H, brm)
Purity	> 90% (NMR)	
MS	626 (M+1)	
Example No.	373	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 11.40 (1H, brs), 9.25 (2H, brs), 8.29 (1H, d, J=1.3Hz), 8.12-8.09 (2H, m), 7.96 (1H, d, J=8.7Hz), 7.88 (1H, dd, J=1.8Hz, 8.1Hz), 7.67-7.63 (2H, m), 7.56 (2H, d, J=8.7Hz), 7.51 (2H, d, J=8.7Hz), 7.17 (1H, d, J=12.0Hz), 7.05 (1H, d, J=8.6Hz), 5.16 (2H, s), 4.05 (1H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	613 (M+1)	

Table 236.

Example No.	374	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.21 (1H, brs), 8.31 (1H, d, J=1.4Hz), 8.18-8.15 (2H, m), 7.99 (1H, d, J=8.7Hz), 7.94 (1H, dd, J=1.8Hz, 8.0Hz), 7.70-7.53 (6H, m), 7.17 (1H, d, J=12.0Hz), 7.05 (1H, d, J=8.6Hz), 5.20 (2H, s), 4.09 (1H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	639 (M+1)	

Example No.	375	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, d, J=1.5Hz), 8.23 (1H, d, J=1.5Hz), 8.19 (1H, d, J=9.0Hz), 8.03-7.98 (2H, m), 7.68 (1H, t, J=8.4Hz), 7.60 (1H, d, J=8.1Hz), 7.56 (2H, d, J=9.3Hz), 7.53 (2H, d, J=9.0Hz), 7.22 (1H, dd, J=2.1Hz, 12.0Hz), 7.09 (1H, dd, J=2.1Hz, 8.4Hz), 5.21 (2H, s), 4.12 (1H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	658 (M+1)	

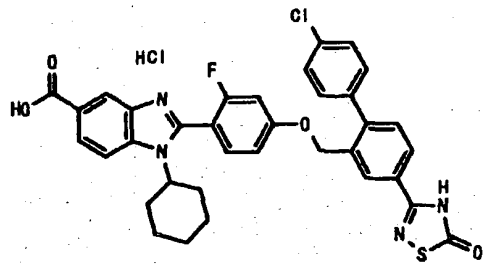
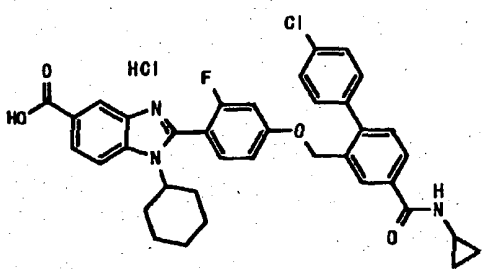
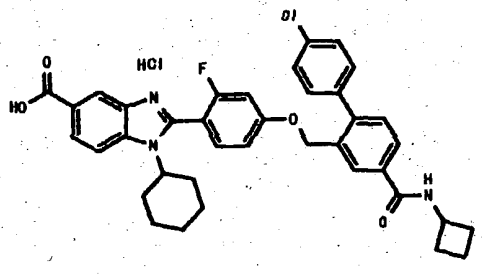
Example No.	376	1H NMR (δ) ppm
		300MHz, DMSO-d6 13.61 (1H, brs), 8.34-8.30 (2H, m), 8.21 (1H, d, J=8.7Hz), 8.07 (1H, dd, J=1.8Hz, 8.1Hz), 8.02 (1H, dd, J=1.5Hz, 8.7Hz), 7.69 (1H, t, J=8.4Hz), 7.57-7.49 (5H, m), 7.22 (1H, dd, J=2.7Hz, 12.0Hz), 7.09 (1H, dd, J=2.4Hz, 9.0Hz), 5.19 (2H, s), 4.12 (1H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	655 (M+1)	

Table 237

Example No.	377	<sup>1</sup> H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.60 (1H, d, J=4.5Hz), 8.29 (1H, d, J=1.5Hz), 8.14 (1H, d, J=8.9Hz), 8.13 (1H, d, J=1.5Hz), 7.98 (1H, dd, J=8.9, 1.5Hz), 7.94 (1H, dd, J=8.1, 1.5Hz), 7.64 (1H, t, J=8.7Hz), 7.52 and 7.49 (4H, ABq, J=9.0Hz), 7.46 (1H, d, J=8.1Hz), 7.18 (1H, dd, J=12.1, 2.3Hz), 7.05 (1H, dd, J=8.7, 2.3Hz), 5.13 (2H, s), 4.08 (1H, brt, J=12.1Hz), 2.95-2.84 (1H, m), 2.31-2.14 (2H, brm), 1.97-1.78 (4H, brm), 1.72-1.59 (1H, brm), 1.47-1.21 (3H, brm), 0.76-0.58 (4H, m)
Purity	> 90% (NMR)	
MS	638 (M+1)	

Example No.	378	<sup>1</sup> H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.77 (1H, d, J=1.4Hz), 8.30 (1H, d, J=1.4Hz), 8.16 (1H, d, J=1.8Hz), 8.13 (1H, d, J=8.4Hz), 7.98 (2H, dd, J=8.4, 1.8Hz), 7.65 (1H, t, J=8.4Hz), 7.53 and 7.49 (4H, ABq, J=8.8Hz), 7.47 (1H, d, J=7.7Hz), 7.18 (1H, dd, J=12.1, 2.2Hz), 7.05 (1H, dd, J=8.4, 2.2Hz), 5.13 (2H, s), 4.53-4.40 (1H, m), 4.09 (1H, brt, J=12.8Hz), 2.31-2.02 (6H, brm), 1.96-1.80 (4H, brm), 1.78-1.60 (3H, brm), 1.47-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	652 (M+1)	

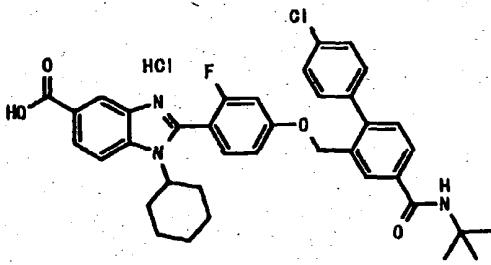
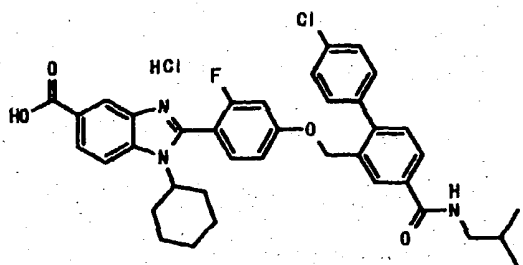
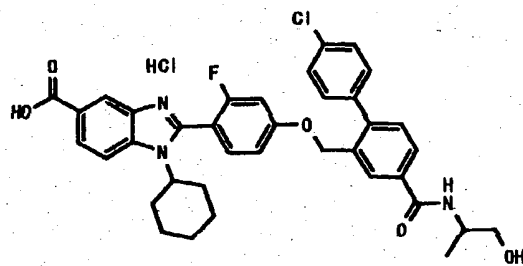
Example No.	379	<sup>1</sup> H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.29 (1H, d, J=1.1Hz), 8.11 (1H, d, J=1.5Hz), 8.11 (1H, d, J=8.8Hz), 7.98-7.91 (2H, m), 7.89 (1H, s), 7.63 (1H, t, J=8.8Hz), 7.52 and 7.48 (4H, ABq, J=8.6Hz), 7.44 (1H, d, J=8.1Hz), 7.17 (1H, dd, J=12.1, 2.2Hz), 7.04 (1H, dd, J=8.8, 2.2Hz), 5.12 (2H, s), 4.07 (1H, brt, J=12.4Hz), 2.33-2.14 (2H, brm), 1.96-1.79 (4H, brm), 1.70-1.60 (1H, brm), 1.48-1.21 (3H, brm), 1.41 (9H, s)
Purity	> 90% (NMR)	
MS	654 (M+1)	



Table 238.

Example No.	380	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.62 (1H, t, J=5.5Hz), 8.30 (1H, d, J=1.5Hz), 8.17 (1H, d, J=1.8Hz), 8.14 (1H, d, J=8.8Hz), 7.98 (1H, dd, J=8.1, 1.8Hz), 7.64 (1H, t, J=8.8Hz), 7.52 and 7.50 (4H, ABq, J=8.8Hz), 7.48 (1H, d, J=8.1Hz), 7.18 (1H, dd, J=12.1, 2.2Hz), 7.05 (1H, dd, J=8.8, 2.2Hz), 5.14 (2H, s), 4.08 (1H, brt, J=12.1Hz), 3.13 (1H, t, J=6.2Hz), 2.31-2.14 (2H, brm), 1.97-1.78 (5H, brm), 1.70-1.60 (1H, brm), 1.47-1.21 (3H, brm), 0.92 (3H, s), 0.90 (3H, s)
Purity	> 90% (NMR)	
MS	654 (M+1)	

Example No.	381	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.29 (1H, d, J=1.5Hz), 8.27 (1H, d, J=8.3Hz), 8.18 (1H, d, J=1.9Hz), 8.13 (1H, d, J=8.7Hz), 8.01-7.96 (2H, m), 7.64 (1H, t, J=8.7Hz), 7.52 and 7.49 (1H, ABq, J=8.8Hz), 7.49 (1H, d, J=7.9Hz), 7.18 (1H, dd, J=12.1, 2.3Hz), 7.05 (1H, dd, J=8.7, 2.3Hz), 5.13 (2H, s), 4.12-4.00 (2H, m), 3.52-3.34 (2H, m), 2.31-2.14 (2H, brm), 1.97-1.79 (4H, brm), 1.71-1.60 (1H, brm), 1.48-1.21 (3H, m), 1.17 and 1.15 (total 3H, each s)
Purity	> 90% (NMR)	
MS	656 (M+1)	

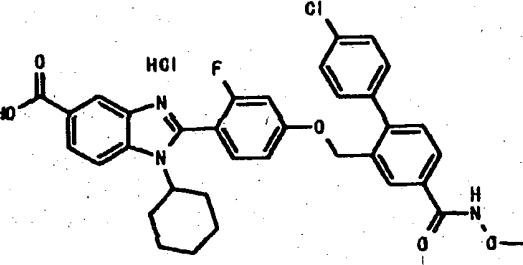
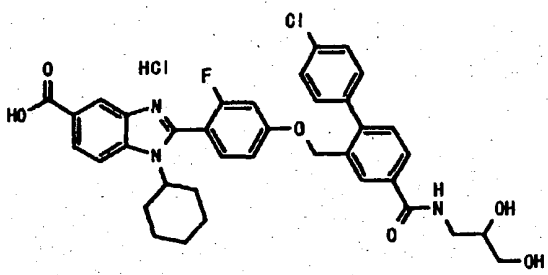
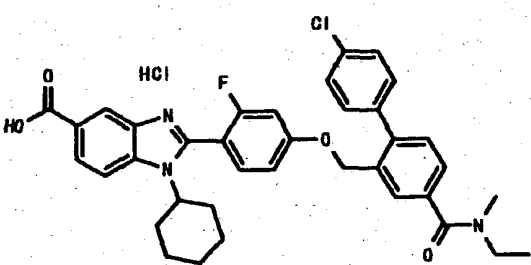
Example No.	382	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.30 (1H, d, J=1.5Hz), 8.13 (1H, d, J=8.8Hz), 8.09 (1H, d, J=1.5Hz), 7.98 (1H, dd, J=8.8, 1.5Hz), 7.86 (1H, dd, J=8.1, 1.5Hz), 7.64 (1H, J=8.8Hz), 7.55-7.47 (5H, m), 7.17 (1H, dd, J=12.1, 2.2Hz), 7.05 (1H, dd, J=8.8, 2.2Hz), 5.14 (2H, s), 4.08 (1H, brt, J=12.8Hz), 3.75 (3H, s), 2.32-2.14 (2H, brm), 1.96-1.78 (4H, brm), 1.70-1.59 (1H, brm), 1.47-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	628 (M+1)	

Table 239

Example No.	383	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.57 (1H, t, J=5.5Hz), 8.29 (1H, d, J=1.4Hz), 8.19 (1H, d, J=1.5Hz), 8.12 (1H, d, J=9.2Hz), 8.01-7.95 (2H, m), 7.64 (1H, t, J=8.8Hz), 7.53 and 7.50 (4H, ABq, J=8.8Hz), 7.48 (1H, d, J=7.7Hz), 7.17 (1H, dd, J=12.1, 2.2Hz), 7.04 (1H, dd, J=8.8, 2.2Hz), 5.14 (2H, s), 4.08 (1H, brt, J=13.9Hz), 3.70-3.66 (1H, m), 3.48-3.36 (3H, m), 3.28-3.20 (1H, m), 2.32-2.13 (2H, brm), 1.96-1.79 (4H, brm), 1.71-1.60 (1H, brm), 1.47-1.19 (3H, brm)
Purity	> 90% (NMR)	
MS	672 (M+1)	

Example No.	384	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.30 (1H, d, J=1.5Hz), 8.14 (1H, d, J=8.4Hz), 7.98 (1H, dd, J=8.4, 1.5Hz), 7.68 (1H, brs), 7.63 (1H, t, J=8.4Hz), 7.51 (5H, s), 7.43 (1H, d, J=8.1Hz), 7.17 (1H, dd, J=12.5, 1.8Hz), 7.03 (1H, dd, J=8.4, 1.8Hz), 4.08 (1H, brt, J=11.4Hz), 3.50 and 3.30 (total 2H, each brs), 2.97 (3H, brs), 2.33-2.13 (2H, brm), 1.96-1.79 (4H, brm), 1.70-1.59 (1H, brm), 1.47-1.03 (6H, brm),
Purity	> 90% (NMR)	
MS	640 (M+1)	

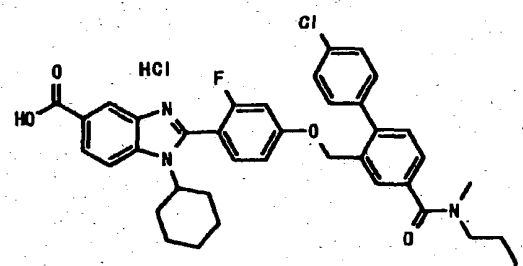
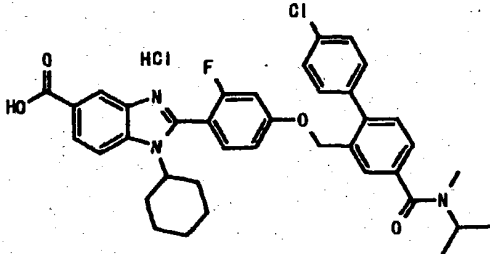
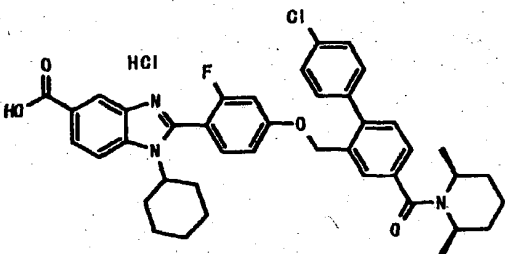
Example No.	385	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.29 (1H, d, J=1.5Hz), 8.12 (1H, d, J=8.8Hz), 7.97 (1H, dd, J=8.8, 1.5Hz), 7.72-7.60 (2H, m), 7.55-7.42 (6H, m), 7.16 (1H, d, J=11.7Hz), 7.03 (1H, d, J=8.4Hz), 5.15 (2H, s), 4.07 (1H, brt, J=12.5Hz), 3.44 and 3.22 (total 2H, each s), 2.97 (3H, brs), 2.32-2.13 (2H, brm), 1.72-1.50 (3H, brm), 1.47-1.23 (3H, brm), 0.93 and 0.72 (total 3H, each brs)
Purity	> 90% (NMR)	
MS	654 (M+1)	

Table 240 -

Example No.	386	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.29 (1H, d, J=1.5Hz), 8.12 (1H, d, J=8.7Hz), 7.97 (1H, dd, J=8.7, 1.5Hz), 7.74-7.60 (2H, m), 7.54-7.42 (6H, m), 7.17 (1H, dd, J=12.1, 2.2Hz), 7.02 (1H, dd, J=8.3, 2.2Hz), 5.15 (2H, s), 4.06 (1H, brt, J=12.8Hz), 3.92 (1H, brs), 2.85 (3H, brs), 2.32-2.14 (2H, brm), 1.96-1.79 (4H, brm), 1.70-1.59 (1H, brm), 1.46-1.07 (3H, brm), 1.15 (6H, brs)
Purity	> 90% (NMR)	
MS	654 (M+1)	

Example No.	387	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.29 (1H, s), 8.14 and 7.97 (2H, ABq, J=8.7Hz), 7.63 (1H, s), 7.63 (1H, t, J=8.7Hz), 7.51-7.41 (6H, m), 7.16 (1H, dd, J=12.1, 1.9Hz), 7.02 (1H, dd, J=8.7, 1.9Hz), 5.16 (2H, s), 4.26 (2H, brs), 4.07 (1H, brt, J=12.1Hz), 2.32-2.14 (2H, brm), 1.97-1.78 (5H, brm), 1.70-1.15 (9H, brm), 1.24 (3H, s), 1.21 (3H, s)
Purity	> 90% (NMR)	
MS	694 (M+1)	

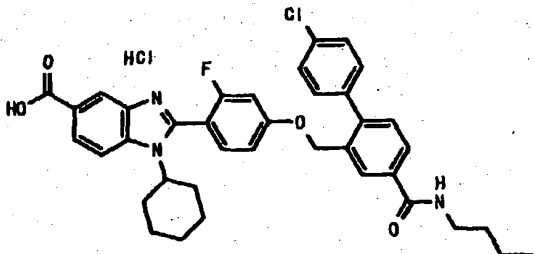
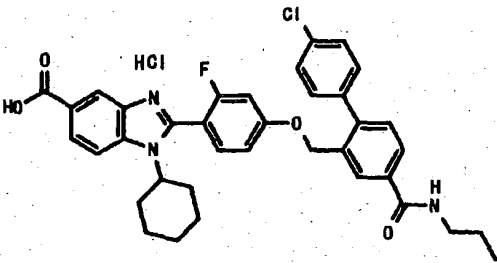
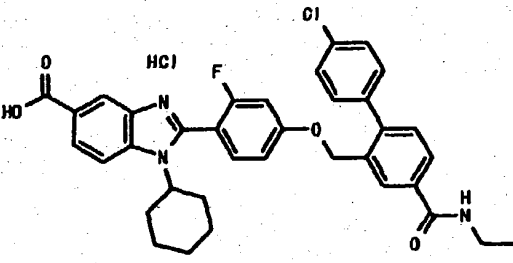
Example No.	388	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.58 (1H, m), 8.29 (1H, s), 8.20-8.10 (2H, m), 8.05-7.90 (2H, m), 7.64 (1H, t, J=8.4Hz), 7.60-7.40 (5H, m), 7.15 (1H, d, J=12.3Hz), 7.04 (1H, d, J=8.4Hz), 5.13 (2H, s), 4.08 (1H, m), 3.40-3.20 (2H, m), 2.35-2.10 (2H, m), 2.00-1.20 (12H, m), 0.91 (3H, t, J=6.9Hz)
Purity	> 90% (NMR)	
MS	654 (M+1)	

Table 241

Example No.	389	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.60 (1H, m), 8.29 (1H, s), 8.20-7.90 (4H, m), 7.64 (1H, t, J=9.0 Hz), 7.60-7.40 (5H, m), 7.17 (1H, d, J=12.0 Hz), 7.04 (1H, d, J=8.7 Hz), 5.13 (2H, s), 4.80 (1H, m), 3.35-3.15 (2H, m), 2.30-2.05 (2H, m), 2.00-1.10 (10H, m), 0.91 (3H, t, J=7.5 Hz)
Purity	> 90% (NMR)	
MS	640 (M+1)	

Example No.	390	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.62 (1H, m), 8.30 (1H, s), 8.20-8.10 (2H, m), 8.05-7.90 (2H, m), 7.65 (1H, t, J=8.4 Hz), 7.60-7.40 (5H, m), 7.18 (1H, d, J=12.0 Hz), 7.05 (1H, d, J=8.4 Hz), 5.14 (2H, s), 4.09 (1H, m), 3.40-3.20 (2H, m), 2.35-2.10 (2H, m), 2.00-1.80 (4H, m), 1.75-1.60 (1H, m), 1.45-1.20 (3H, m), 1.15 (3H, t, J=7.2 Hz)
Purity	> 90% (NMR)	
MS	626 (M+1)	

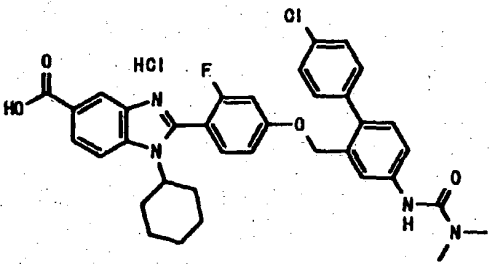
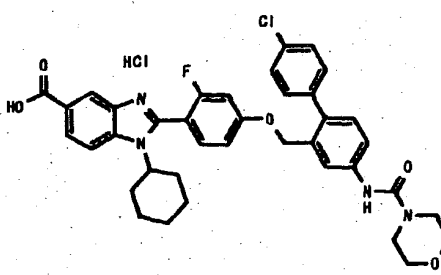
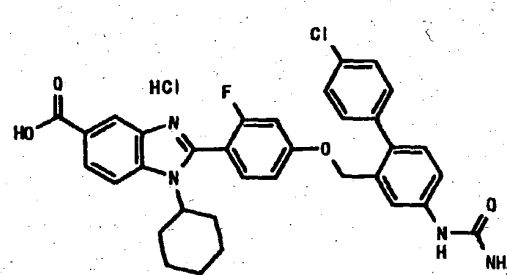
Example No.	391	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.54 (1H, s), 8.31 (1H, s), 8.19 (1H, d, J=8.6 Hz), 8.01 (1H, d, J=8.6 Hz), 7.81 (1H, d, J=2.1 Hz), 7.64 (1H, t, J=8.4 Hz), 7.61 (1H, dd, J=2.3 Hz, 8.4 Hz), 7.47 (2H, d, J=8.6 Hz), 7.43 (2H, d, J=8.8 Hz), 7.25 (1H, d, J=8.4 Hz), 7.17 (1H, dd, J=2.3 Hz, 12.1 Hz), 7.05 (1H, dd, J=2.3 Hz, 8.6 Hz), 5.05 (2H, s), 4.12 (1H, m), 2.96 (6H, s), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	641 (M+1)	

Table 242

Example No.	392	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.79(1H, s), 8.29(1H, d, J=1.5Hz), 8.13(1H, d, J=8.8Hz), 7.98(1H, dd, J=8.8, 1.5Hz), 7.80(1H, d, J=2.2Hz), 7.63(1H, t, J=8.4Hz), 7.61(1H, dd, J=8.2, 2.2Hz), 7.47 and 7.43(4H, ABq, J=8.8Hz), 7.26(1H, d, J=8.2Hz), 7.14(1H, dd, J=12.1, 2.2Hz), 7.02(1H, d, J=8.4, 2.2Hz), 5.05(2H, s), 4.08(1H, brt, J=12.1Hz), 3.64-3.61(2H, m), 3.48-3.45(2H, m), 2.32-2.13(2H, brm), 1.96-1.78(4H, brm), 1.70-1.66(1H, brm), 1.44-1.19(3H, brm)
Purity	> 90% (NMR)	
MS	683(M+1)	

Example No.	393	<sup>1</sup> H NMR (δ) ppm
		400MHz, DMSO-d <sub>6</sub> 8.94(1H, s), 8.31(1H, d, J=1.0Hz), 8.18(1H, d, J=8.6Hz), 8.00(1H, dd, J=1.4Hz, 8.8Hz), 7.71(1H, d, J=2.2Hz), 7.66(1H, t, J=8.6Hz), 7.52(1H, dd, J=2.4Hz, 8.6Hz), 7.46(2H, d, J=8.6Hz), 7.42(2H, d, J=8.2Hz), 7.24(1H, d, J=8.4Hz), 7.16(1H, d, J=12.1Hz), 7.04(1H, dd, J=2.4Hz, 8.8Hz), 5.05(2H, s), 4.13(1H, m), 2.40-2.10(2H, m), 2.00-1.75(4H, m), 1.70-1.55(1H, m), 1.50-1.20(3H, m)
Purity	> 90% (NMR)	
MS	613(M+1)	

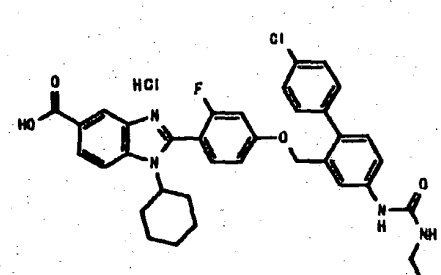
Example No.	394	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.93(1H, s), 8.31(1H, d, J=1.4Hz), 8.19(1H, d, J=8.8Hz), 8.01(1H, d, J=8.7Hz), 7.71(1H, d, J=2.2Hz), 7.66(1H, t, J=8.5Hz), 7.51(1H, dd, J=2.2Hz, 8.4Hz), 7.46(2H, d, J=8.6Hz), 7.41(2H, d, J=8.7Hz), 7.23(1H, d, J=8.4Hz), 7.16(1H, d, J=12.2Hz), 7.05(1H, d, J=8.7Hz), 5.05(2H, s), 4.13(1H, m), 3.12(2H, q, J=7.2Hz), 2.40-2.10(2H, m), 2.00-1.75(4H, m), 1.70-1.60(1H, m), 1.55-1.20(3H, m), 1.06(3H, t, J=7.2Hz)
Purity	> 90% (NMR)	
MS	641(M+1)	

Table 243

Example No.	395	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.83 (1H, s), 8.32 (1H, d, J=1.4 Hz), 8.21 (1H, d, J=8.8 Hz), 8.02 (1H, dd, J=1.4 Hz, 8.7 Hz), 7.71 (1H, d, J=2.1 Hz), 7.68 (1H, t, J=8.6 Hz), 7.49 (1H, dd, J=2.2 Hz, 8.4 Hz), 7.46 (2H, d, J=8.4 Hz), 7.41 (2H, d, J=8.6 Hz), 7.23 (1H, d, J=8.4 Hz), 7.17 (1H, d, J=12.2 Hz), 7.06 (1H, d, J=8.7 Hz), 6.30 (1H, brs), 5.05 (2H, s), 4.14 (1H, m), 3.77 (1H, sept, J=6.5 Hz), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m), 1.11 (6H, d, J=6.5 Hz)
Purity	> 90% (NMR)	
MS	655 (M+1)	

Example No.	396	<sup>1</sup> H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.37 (1H, d, J=7.3Hz), 8.25 (1H, s), 8.15 (1H, s), 7.97 (2H, d, J=8.8Hz), 7.88 (1H, d, J=8.8Hz), 7.58-7.47 (4H, m), 7.31 (1H, m), 7.11 (1H, dd, J=8.4, 2.2Hz), 6.98 (1H, dd, J=8.4, 2.2), 5.13 (2H, s), 4.13 (1H, q, J=6.6Hz), 3.98 (1H, m), 2.19 (2H, m), 1.86 (4H, m) 1.62 (1H, m) 1.31 (3H, m), 1.20 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	642 (M+1)	

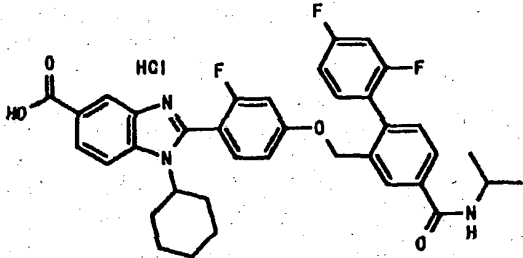
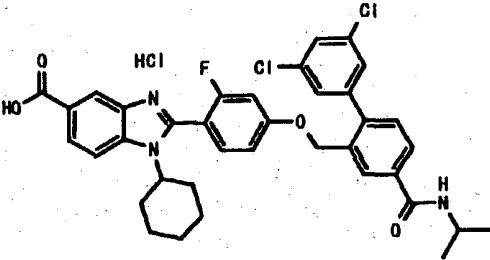
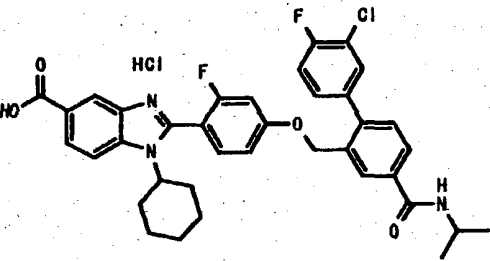
Example No.	397	<sup>1</sup> H NMR (δ) ppm 300MHz, DMSO-d <sub>6</sub> 8.40 (1H, d, J=7.9Hz), 8.28 (1H, d, J=1.9Hz), 8.15 (1H, d, J=1.9Hz), 8.11 (1H, d, J=8.7Hz), 7.96 (2H, m), 7.56 (1H, t, J=8.7Hz), 7.45 (3H, m), 7.18 (1H, m), 7.08 (1H, dd, J=12.1, 1.9Hz), 6.96 (1H, d, J=8.3, 2.3Hz), 5.09 (2H, s), 4.14 (1H, m), 4.04 (1H, m), 2.23 (2H, m), 1.86 (3H, m), 1.62 (1H, m), 1.33 (3H, m), 1.20 (6H, d, J=6.4Hz)
		
Purity	> 90% (NMR)	
MS	642 (M+1)	

Table 244

Example No.	398	1H NMR (δ) ppm
		8.41 (1H, d, J=8.1Hz), 8.29 (1H, d, J=1.5Hz), 8.17 (1H, d, J=1.8Hz), 8.12 (1H, d, J=8.4Hz), 8.01-7.95 (2H, m), 7.67-7.62 (2H, m), 7.55-7.51 (3H, m), 7.19 (1H, dd, J=12.1, 2.2Hz), 7.05 (1H, dd, J=8.8, 2.2Hz), 5.13 (2H, s), 4.10-4.00 (2H, m), 2.32-2.13 (4H, m), 1.71-1.60 (1H, m), 1.49-1.14 (3H, m), 1.21 (3H, s), 1.19 (3H, s)
Purity	> 90% (NMR)	
MS	674 (M+1)	

Example No.	399	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.39 (1H, d, J=7.7Hz), 8.29 (1H, d, J=1.5Hz), 8.16 (1H, d, J=1.8Hz), 8.11 (1H, d, J=8.8Hz), 8.00-7.95 (2H, m), 7.69-7.61 (2H, m), 7.54-7.46 (3H, m), 7.18 (1H, dd, J=12.1, 2.2Hz), 7.04 (1H, dd, J=8.8, 2.2Hz), 5.13 (2H, s), 4.20-4.02 (2H, m), 2.33-2.13 (2H, brm), 1.97-1.80 (4H, m), 1.72-1.61 (1H, m), 1.44-1.13 (3H, m), 1.21 (3H, s), 1.19 (3H, s)
Purity	> 90% (NMR)	
MS	658 (M+1)	

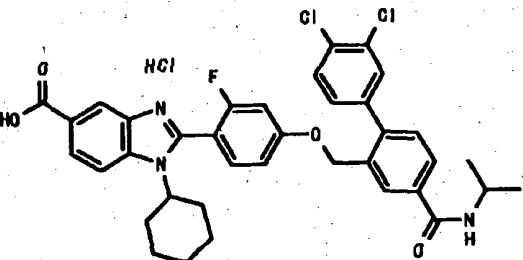
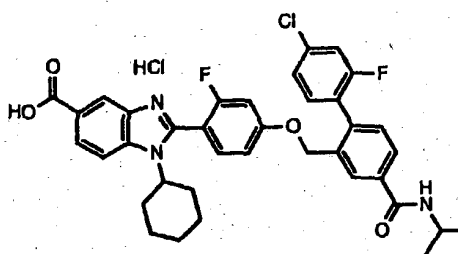
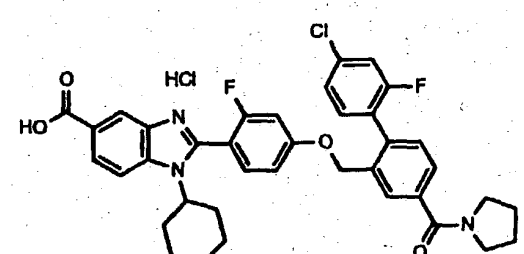
Example No.	400	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.39 (1H, d, J=7.7Hz), 8.29 (1H, s), 8.17 (1H, d, J=1.5Hz), 8.11 (1H, d, J=8.8Hz), 7.98 (2H, m), 7.73 (2H, m), 7.64 (1H, t, J=8.4Hz), 7.52 (1H, d, J=8.0Hz), 7.46 (1H, dd, J=8.4, 1.8Hz), 7.18 (1H, dd, J=11.9, 2.0Hz), 7.05 (1H, dd, J=8.6, 2.4Hz), 5.14 (2H, s), 4.13 (2H, m), 2.22 (2H, m), 1.88 (4H, m), 1.64 (1H, m), 1.34 (3H, m), 1.20 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	642 (M+1)	

Table 245

Example No.	401	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.38 (1H, d, J=7.8Hz), 8.28 (1H, s), 8.20-8.05 (2H, m), 8.00-7.90 (2H, m), 7.65-7.30 (5H, m), 7.09 (1H, d, J=12.3Hz), 6.97 (1H, d, J=10.2Hz), 5.09 (2H, s), 4.20-4.00 (2H, m), 2.30-2.10 (2H, m), 2.00-1.80 (4H, m), 1.70-1.60 (1H, m), 1.40-1.10 (3H, m), 1.19 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	658 (M+1)	

Example No.	402	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.25 (1H, s), 8.03 (1H, d, J=8.7Hz), 7.91 (1H, d, J=8.7Hz), 7.83 (1H, s), 7.70-7.35 (6H, m), 7.04 (1H, d, J=12.0Hz), 6.93 (1H, d, J=8.4Hz), 5.09 (2H, s), 4.00 (1H, m), 3.60-3.40 (4H, m), 2.30-2.10 (2H, m), 1.45-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	670 (M+1)	

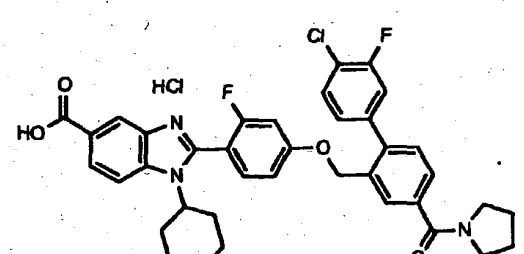
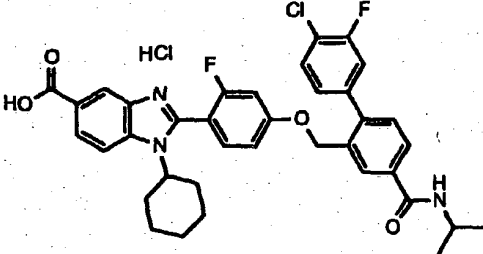
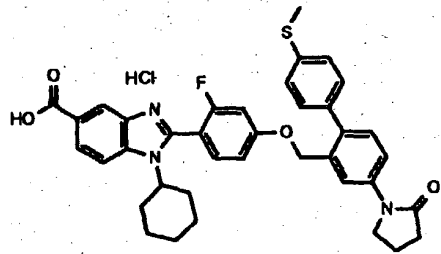
Example No.	403	1H NMR (δ) ppm
		400MHz, DMSO-d <sub>6</sub> 8.25 (1H, s), 8.08 (1H, d, J=8.4Hz), 7.92 (1H, d, J=9.2Hz), 7.79 (1H, s), 7.66-7.49 (4H, m), 7.42 (1H, d, J=7.6Hz), 7.31-7.28 (1H, m), 7.14 (1H, d, J=11.3Hz), 6.99 (1H, d, J=8.8Hz), 5.13 (2H, s), 4.02 (1H, m), 3.54-3.33 (4H, m), 2.29-2.08 (2H, m), 1.93-1.73 (8H, m), 1.67-1.52 (1H, m), 1.48-1.11 (3H, m)
Purity	> 90% (NMR)	
MS	670 (M+1)	



Table 246

Example No.	404	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.41 (1H, d, J=7.6Hz), 8.32 (1H, d, J=1.5Hz), 8.20 (1H, d, J=8.6Hz), 8.17 (1H, d, J=1.7Hz), 8.00 (1H, dt, J=8.8Hz, 1.5Hz), 7.71-7.64 (2H, m), 7.54 (1H, dd, J=10.3Hz, 1.9Hz), 7.32 (1H, dd, J=8.2Hz, 1.9Hz), 7.22 (1H, dd, J=12.1Hz, 2.3Hz), 7.08 (1H, dd, J=8.6Hz, 2.3Hz), 5.17 (2H, s), 4.15 (1H, m), 2.31-2.14 (2H, m), 1.99-1.70 (4H, m), 1.70-1.60 (1H, m), 1.46-1.20 (3H, m), 1.19 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	658 (M+1)	

Example No.	405	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.32 (1H, s), 8.19 (1H, d, J=9.0Hz), 8.03-7.98 (2H, m), 7.75 (1H, dd, J=2.1Hz, 8.4Hz), 7.67 (1H, t, J=8.6Hz), 7.40-7.36 (3H, m), 7.32 (2H, d, J=8.4Hz), 7.19 (1H, d, J=2.1Hz, 12.3Hz), 7.07 (1H, d, J=2.1Hz, 8.7Hz), 5.11 (2H, s), 4.12 (1H, m), 4.12 (1H, m), 3.90 (2H, t, J=6.9Hz), 2.54 (2H, t, J=8.1Hz), 2.50 (3H, s), 2.40-2.05 (4H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	650 (M+1)	

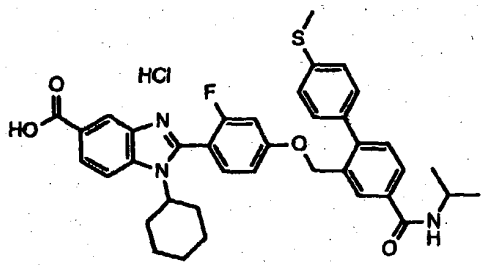
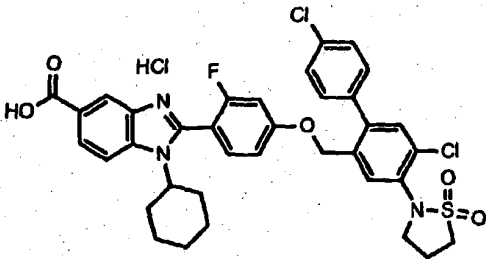
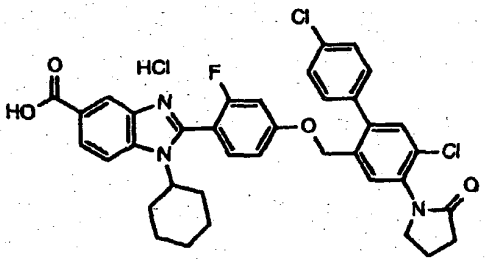
Example No.	406	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.34 (1H, d, J=7.7Hz), 8.29 (1H, s), 8.15 (1H, s), 8.11 (1H, d, J=8.8Hz), 7.97 (2H, d, J=9.2Hz), 7.63 (1H, t, J=8.8Hz), 7.47-7.31 (5H, m), 7.18 (1H, dd, J=12.4, 2.2Hz), 7.06 (1H, dd, J=12.4, 2.2Hz), 5.13 (2H, s), 4.13 (2H, m), 1.96 (2H, m), 1.87 (4H, m), 1.62 (1H, m), 1.34 (3H, m), 1.20 (6H, d, J=6.2Hz)
Purity	> 90% (NMR)	
MS	652 (M+1)	

Table 247-

Example No.	407	1H NMR(δ) ppm
		400MHz, DMSO-d <sub>6</sub> 8.32(1H, d, J=1.4Hz), 8.20(1H, d, J=8.8Hz), 8.01(1H, dd, J=1.6Hz, 8.8Hz), 7.90(1H, s), 7.67(1H, t, J=8.4Hz), 7.61(1H, s), 7.55-7.50(4H, m), 7.21(1H, dd, J=2.3Hz, 12.0Hz), 7.06(1H, dd, J=2.2Hz, 8.7Hz), 5.10(2H, s), 4.11(1H, m), 3.78(2H, t, J=6.7Hz), 3.47(2H, t, J=7.4Hz), 2.54-2.48(2H, m), 2.40-2.10(2H, m), 2.00-1.80(4H, m), 1.75-1.55(1H, m), 1.50-1.20(3H, m)
Purity	> 90% (NMR)	
MS	708(M+1)	

Example No.	408	1H NMR(δ) ppm
		400MHz, DMSO-d <sub>6</sub> 8.32(1H, d, J=1.6Hz), 8.21(1H, d, J=8.8Hz), 8.02(1H, dd, J=1.6Hz, 8.8Hz), 7.76(1H, s), 7.68(1H, t, J=8.5Hz), 7.59(1H, s), 7.54-7.51(4H, m), 7.21(1H, dd, J=2.4Hz, 12.1Hz), 7.07(1H, dd, J=2.4Hz, 8.8Hz), 5.08(2H, s), 4.11(1H, m), 3.77(2H, t, J=6.9Hz), 2.47(2H, t, J=8.0Hz), 2.40-2.10(4H, m), 2.00-1.80(4H, m), 1.70-1.60(1H, m), 1.45-1.20(3H, m)
Purity	> 90% (NMR)	
MS	672(M+1)	

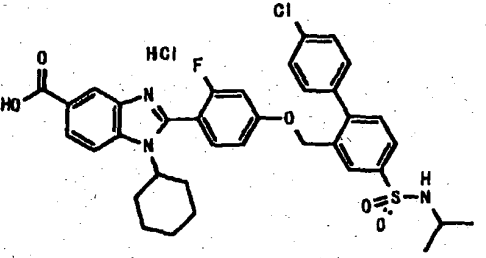
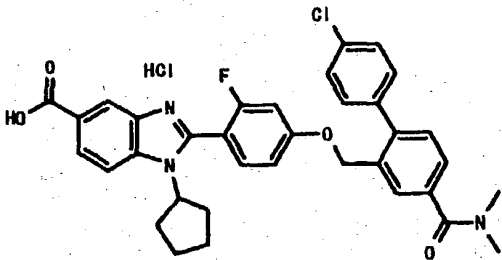
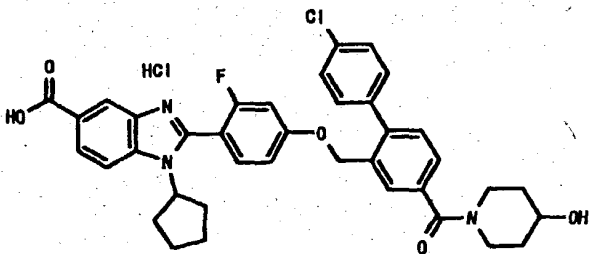
Example No.	409	1H NMR(δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.28(1H, d, J=1.5Hz), 8.20-8.85(4H, m), 7.75(1H, d, J=6.9Hz), 7.70-7.45(6H, m), 7.13(1H, dd, J=12.0Hz, 2.1Hz), 7.00(1H, dd, J=8.7Hz), 2.1Hz), 5.22(2H, s), 4.05(1H, m), 3.40-3.20(1H, m), 2.30-2.10(2H, m), 2.00-1.55(5H, m), 1.45-1.10(3H, m), 1.00(6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	676(M+1)	

Table 248

Example No.	410	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.31 (1H, s), 8.00 (1H, d, J=8.7Hz), 7.88 (1H, d, J=8.7Hz), 7.70 (1H, s), 7.65 (1H, t, J=8.4Hz), 7.53 (2H, d, J=8.4Hz), 7.49 (2H, d, J=8.7Hz), 7.45-7.41 (2H, m), 7.16 (1H, d, J=12.0Hz), 7.04 (1H, d, J=8.7Hz), 5.14 (2H, s), 4.68 (1H, quint, J=8.4Hz), 3.02, 2.98 (6H, s), 2.30-1.85 (6H, m), 1.80-1.50 (2H, m)
Purity	> 90% (NMR)	
MS	612 (M+1)	

Example No.	411	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.30 (1H, s), 7.99 (1H, d, J=9.0Hz), 7.87 (1H, d, J=8.7Hz), 7.67 (1H, s), 7.64 (1H, t, J=8.7Hz), 7.53 (2H, d, J=8.7Hz), 7.49 (2H, d, J=7.5Hz), 7.45-7.41 (2H, m), 7.15 (1H, d, J=12.3Hz), 7.02 (1H, d, J=8.4Hz), 5.15 (2H, s), 4.67 (1H, quint, J=8.7Hz), 4.02 (1H, m), 3.76 (1H, m), 3.55 (1H, m), 3.22 (2H, m), 2.40-1.20 (12H, m)
Purity	> 90% (NMR)	
MS	668 (M+1)	

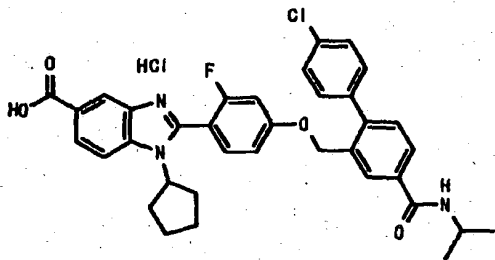
Example No.	412	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.38 (1H, d, J=7.5Hz), 8.33 (1H, s), 8.16 (1H, s), 8.02 (1H, d, J=8.7Hz), 7.98 (1H, d, J=9.0Hz), 7.91 (1H, d, J=8.4Hz), 7.67 (1H, t, J=8.4Hz), 7.53 (2H, d, J=8.7Hz), 7.48 (2H, d, J=8.7Hz), 7.46 (1H, d, J=8.1Hz), 7.18 (1H, d, J=11.7Hz), 7.06 (1H, d, J=8.7Hz), 5.13 (2H, s), 4.70 (1H, quint, J=8.4Hz), 4.13 (1H, sept, J=6.6Hz), 2.30-1.85 (6H, m), 1.80-1.50 (2H, m), 1.16 (6H, d, J=6.3Hz)
Purity	> 90% (NMR)	
MS	626 (M+1)	

Table 249

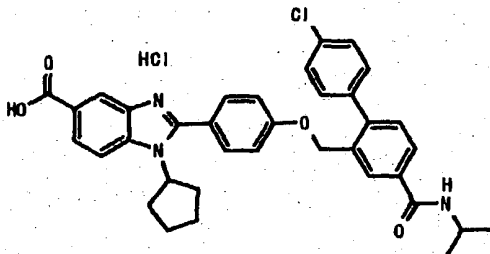
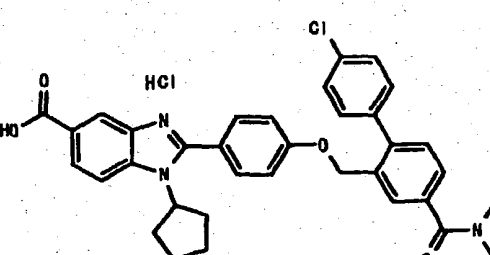
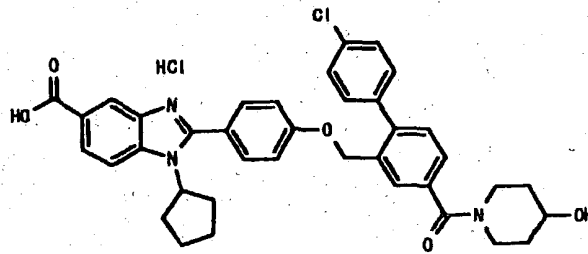
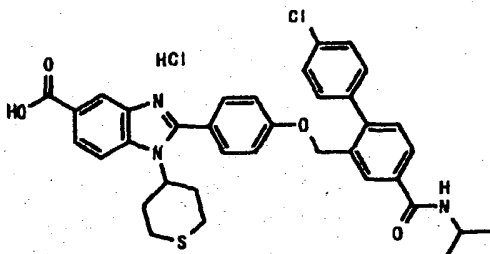
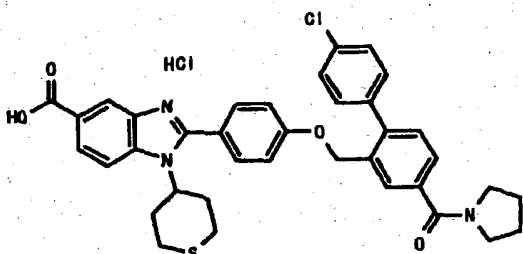
Example No.	413	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.39 (1H, d, J=7.5Hz), 8.31 (1H, d, J=1.5Hz), 8.16 (1H, d, J=1.9Hz), 8.06 (1H, dd, J=8.8, 1.5Hz), 7.99-7.95 (2H, m), 7.76 and 7.24 (4H, ABq, J=8.9Hz), 7.53 and 7.50 (4H, A'B'q, J=9.1Hz), 7.46 (1H, d, J=8.3Hz), 5.14 (2H, s), 4.94 (1H, quint, J=9.0Hz), 4.19-4.08 (1H, m), 2.32-2.11 (4H, brm), 2.10-1.95 (2H, brm), 1.78-1.62 (2H, brm), 1.26 (3H, s), 1.18 (3H, s)
Purity	> 90% (NMR)	
MS	608 (M+1)	
Example No.	414	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.31 (1H, d, J=1.5Hz), 8.06 (1H, dd, J=8.7, 1.5Hz), 7.97 (1H, d, J=8.7Hz), 7.75 and 7.22 (4H, ABq, J=8.9Hz), 7.70 (1H, d, J=1.9Hz), 7.53 (1H, dd, J=7.9, 1.9Hz), 7.52 (4H, s), 7.43 (1H, d, J=7.9Hz), 5.15 (2H, s), 4.93 (1H, quint, J=8.9Hz), 3.01 (3H, s), 2.97 (3H, s), 2.32-2.11 (4H, brm), 2.09-1.94 (2H, brm), 1.77-1.62 (2H, brm)
Purity	> 90% (NMR)	
MS	594 (M+1)	
Example No.	415	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.31 (1H, d, J=1.5Hz), 8.06 (1H, dd, J=8.7, 1.5Hz), 7.98 (1H, d, J=8.7Hz), 7.75 and 7.22 (4H, ABq, J=8.9Hz), 7.67 (1H, d, J=1.5Hz), 7.52 (4H, s), 7.49 (1H, dd, J=7.9, 1.5Hz), 7.43 (1H, d, J=8.9Hz), 5.16 (2H, s), 4.93 (1H, quint, J=8.9Hz), 3.76 (1H, brs), 3.55 (2H, brs), 3.22 (2H, brs), 2.31-2.11 (4H, brm), 2.16-1.95 (2H, brm), 1.88-1.62 (4H, brm), 1.48-1.28 (2H, brm)
Purity	> 90% (NMR)	
MS	650 (M+1)	

Table 250

Example No.	416	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.38 (1H, d, J=7.7Hz), 8.30 (1H, s), 8.20-7.90 (4H, m), 7.72 (2H, d, J=8.7Hz), 7.60-7.40 (5H, m), 7.22 (2H, d, J=8.7Hz), 5.13 (2H, s), 4.47 (1H, m), 4.15 (1H, m), 2.90-2.70 (4H, m), 2.60-2.30 (4H, m), 1.19 (6H, d, J=6.5Hz)
Purity	> 90% (NMR)	
MS	640 (M+1)	

Example No.	417	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.33 (1H, s), 8.17 (1H, d, J=8.6Hz), 8.10 (1H, d, J=8.6Hz), 7.82 (1H, d, J=1.4Hz), 7.74 (2H, d, J=8.7Hz), 7.64 (1H, dd, J=8.0Hz, 1.7Hz), 7.55-7.50 (4H, m), 7.43 (1H, d, J=7.8Hz), 7.24 (1H, d, J=8.7Hz), 5.16 (2H, s), 4.49 (1H, m), 3.60-3.40 (4H, m), 2.90-2.70 (4H, m), 2.60-2.30 (4H, m), 2.20-1.80 (4H, m)
Purity	> 90% (NMR)	
MS	652 (M+1)	

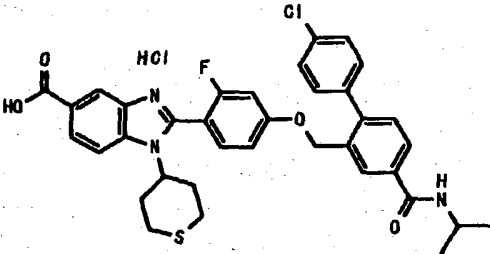
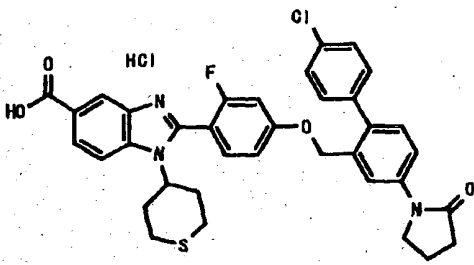
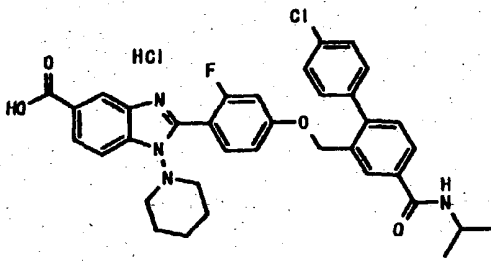
Example No.	418	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.34 (1H, d, J=7.6Hz), 8.25 (1H, s), 8.11 (1H, d, J=1.3Hz), 7.90-8.00 (3H, m), 7.59 (1H, t, J=8.6Hz), 7.40-7.55 (5H, m), 7.12 (1H, d, J=11.9Hz), 7.00 (1H, d, J=8.6Hz), 5.08 (2H, s), 4.30-4.10 (2H, m), 2.80-2.65 (4H, m), 2.45-2.30 (2H, m), 1.15 (6H, d, J=4.8Hz)
Purity	> 90% (NMR)	
MS	658 (M+1)	

Table 251

Example No.	419	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.30 (1H, s), 8.05-7.95 (3H, m), 7.80-7.75 (1H, m), 7.63 (1H, t, J =8.6Hz), 7.55-7.35 (5H, m), 7.1 5 (1H, dd, J=12.1Hz, 2.1Hz), 7.0 3 (1H, dd, J=8.7Hz, 2.3Hz), 5.10 (2H, s), 4.23 (1H, m), 3.90 (2H, t , J=7.0Hz), 2.95-2.70 (4H, m), 2 .60-2.35 (4H, m), 2.30-2.00 (4H , m)
Purity	> 90 % (NMR)	
MS	656 (M+1)	

Example No.	420	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.37 (1H, d, J=7.5Hz), 8.28 (1H, d, J=1.5Hz), 8.17 (1H, d, J=1.5H z), 8.13 (1H, d, J=8.7Hz), 7.97 ( 1H, dd, J=8.1, 1.5Hz), 7.94 (1H, dd, J=8.7, 1.5Hz), 7.61 (1H, t, J =8.7Hz), 7.51 and 7.49 (4H, ABq, J=8.9Hz), 7.46 (1H, d, J=8.1Hz) , 7.08 (1H, dd, J=12.4, 2.3Hz), 6 .97 (1H, dd, J=8.7, 2.3Hz), 5.10 (2H, s), 4.20-4.08 (1H, m), 3.62 -3.56 (2H, brm), 3.13-3.10 (2H, brm), 1.79-1.60 (3H, brm), 1.54 -1.34 (3H, brm), 1.21 (3H, s), 1. 18 (3H, s)
Purity	> 90 % (NMR)	
MS	641 (M+1)	

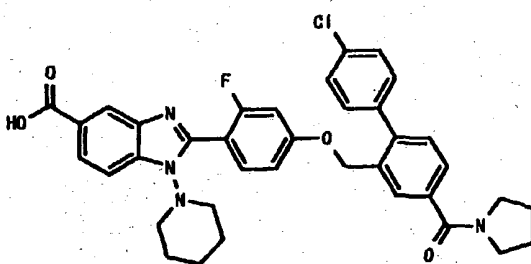
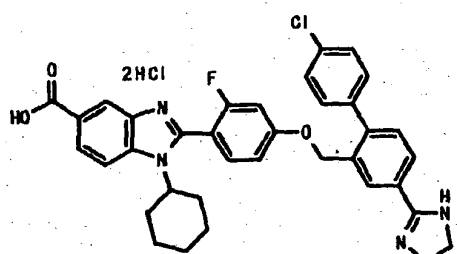
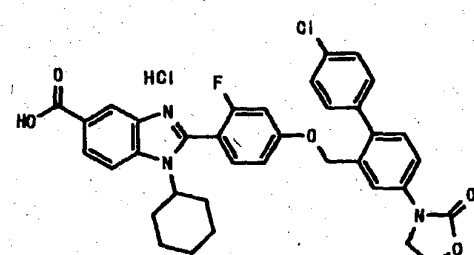
Example No.	421	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.24 (1H, d, J=1.5Hz), 8.02 (1H, d, J=8.7Hz), 7.88 (1H, dd, J=8.7 , 1.5Hz), 7.82 (1H, d, J=1.9Hz), 7.63 (1H, dd, J=7.9, 1.9Hz), 7.5 4 (1H, t, J=8.7Hz), 7.50 (4H, s), 7.42 (1H, d, J=7.9Hz), 7.01 (1H, dd, J=12.0, 2.3Hz), 6.91 (1H, dd , J=8.7, 2.3Hz), 5.11 (2H, s), 3. 63-3.41 (6H, m), 3.07-3.04 (2H, brm), 1.95-1.79 (4H, brm), 1.77 -1.57 (3H, brm), 1.50-1.32 (3H, brm)
Purity	> 90 % (NMR)	
MS	653 (M+1)	

Table 252

Example No.	422	1H NMR (δ) ppm
		300MHz, DMSO-d6 10.99 (2H, s), 8.44 (1H, s), 8.30 (1H, s), 8.18 (1H, d, J=8.7Hz), 8.14 (1H, d, J=8.7Hz), 7.98 (1H, d, J=9.0Hz), 7.70-7.66 (2H, m), 7.57 (2H, d, J=8.7Hz), 7.54 (2H, d, J=8.7Hz), 7.21 (1H, d, J=12.0Hz), 7.09 (1H, d, J=8.4Hz), 5.19 (2H, s), 4.05 (4H, s), 2.40-2.18 (2H, m), 2.15-1.80 (4H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90 % (NMR)	
MS	623 (M+1)	

Example No.	423	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.27 (1H, s), 8.05 (1H, d, J=8.7Hz), 7.93 (1H, d, J=8.7Hz), 7.90 (1H, s), 7.70 (1H, d, J=8.4Hz), 7.59 (1H, t, J=8.4Hz), 7.50 (2H, d, J=9.0Hz), 7.45 (2H, d, J=8.7Hz), 7.41 (1H, d, J=8.4Hz), 7.12 (1H, d, J=12.0Hz), 7.00 (1H, d, J=8.7Hz), 5.10 (2H, s), 4.49 (2H, t, J=7.8Hz), 4.14 (2H, t, J=8.0Hz), 4.04 (1H, m), 2.40-2.10 (2H, m), 2.00-1.50 (5H, m), 1.45-1.20 (3H, m)
Purity	> 90 % (NMR)	
MS	640 (M+1)	

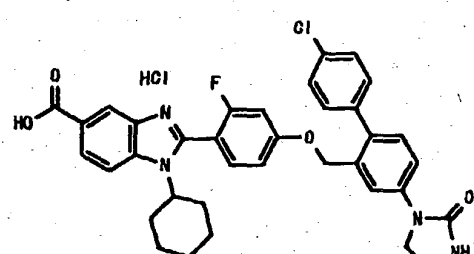
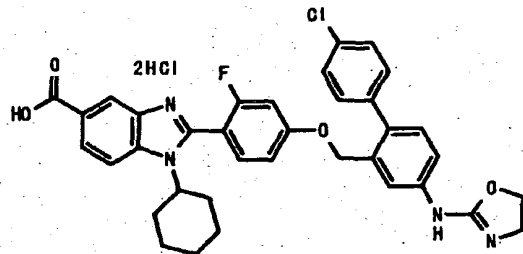
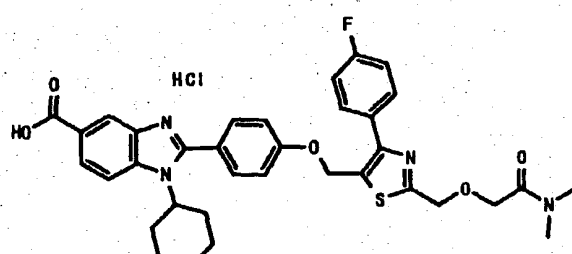
Example No.	424	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.30 (1H, s), 8.14 (1H, d, J=8.4Hz), 7.98 (1H, d, J=9.3Hz), 7.89 (1H, s), 7.68 (1H, d, J=8.4Hz), 7.62 (1H, d, J=9.0Hz), 7.48 (2H, d, J=8.4Hz), 7.43 (2H, d, J=8.4Hz), 7.33 (1H, d, J=8.4Hz), 7.16 (1H, d, J=12.0Hz), 7.04 (1H, d, J=9.0Hz), 5.07 (2H, s), 4.10 (1H, m), 3.92 (2H, t, J=8.0Hz), 3.45 (2H, t, J=8.0Hz), 2.40-2.10 (2H, m), 2.00-1.50 (5H, m), 1.45-1.20 (3H, m)
Purity	> 90 % (NMR)	
MS	639 (M+1)	

Table 253

Example No.	425	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 9.05 (1H, s), 8.30 (1H, s), 8.16 (1H, d, J=8.8Hz), 7.99 (1H, d, J=8.6Hz), 7.72 (1H, s), 7.64 (1H, t, J=8.6Hz), 7.52 (1H, d, J=8.4Hz), 7.47 (2H, d, J=8.7Hz), 7.42 (2H, d, J=8.6Hz), 7.25 (1H, d, J=8.4Hz), 7.15 (1H, d, J=12.2Hz), 7.04 (1H, d, J=8.6Hz), 6.60 (1H, brs), 5.05 (2H, s), 4.10 (1H, m), 3.68 (2H, t, J=6.1Hz), 3.45 (2H, t, J=6.1Hz), 2.40-2.10 (2H, m), 2.00-1.55 (5H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	639 (M+1)	

Example No.	426	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.32 (1H, s), 8.24 (1H, d, J=8.7Hz), 8.03 (1H, d, J=8.7Hz), 7.78-7.73 (4H, m), 7.38-7.32 (4H, m), 5.52 (2H, s), 4.88 (2H, s), 4.40 (2H, s), 4.37 (1H, m), 2.92, 2.84 (6H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	643 (M+1)	

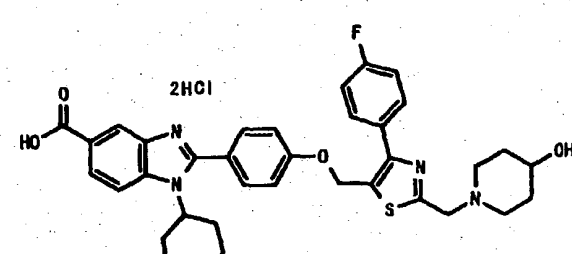
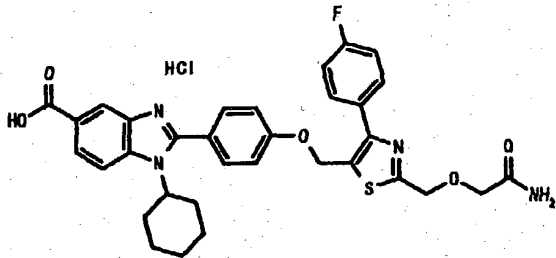
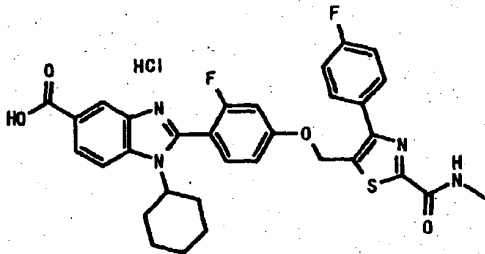
Example No.	427	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 11.26 (1H, brs), 8.35 (1H, s), 8.27 (1H, d, J=9.0Hz), 8.05 (1H, d, J=8.4Hz), 7.83-7.78 (4H, m), 7.42-7.35 (4H, m), 5.57 (2H, s), 4.77, 4.73 (2H, s), 4.37 (1H, m), 3.95 (1H, s), 3.70-3.00 (4H, m), 2.40-1.00 (14H, m)
Purity	> 90% (NMR)	
MS	641 (M+1)	



Table 254

Example No.	428	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.26 (1H, d, J=9.0 Hz), 8.04 (1H, d, J=8.7 Hz), 7.79-7.73 (4H, m), 7.38-7.31 (6H, m), 5.53 (2H, s), 4.90 (2H, s), 4.37 (1H, m), 4.05 (2H, s), 2.40-2.18 (2H, m), 2.15-1.95 (2H, m), 1.90-1.80 (2H, m), 1.75-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	615 (M+1)	

Example No.	429	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.88 (1H, q, J=4.5 Hz), 8.33 (1H, d, J=1.5 Hz), 8.18 (1H, d, J=8.7 Hz), 8.01 (1H, dd, J=1.5 Hz, 8.7 Hz), 7.89-7.83 (2H, m), 7.50-7.34 (3H, m), 7.20 (1H, dd, J=2.1 Hz, 8.4 Hz), 5.61 (2H, s), 4.13 (1H, m), 2.84 (3H, d, J=4.8 Hz), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	603 (M+1)	

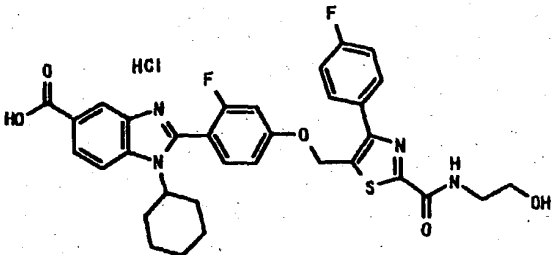
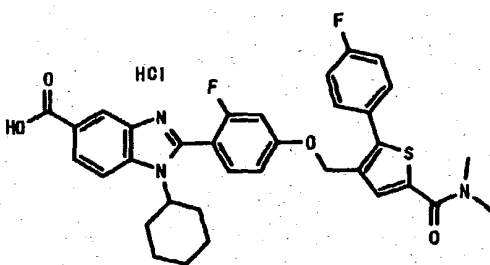
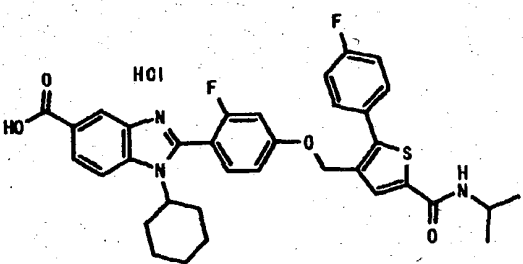
Example No.	430	1H NMR (δ) ppm
		400MHz, DMSO-d6 8.79 (1H, t, J=5.9 Hz), 8.31 (1H, s), 8.15 (1H, d, J=8.7 Hz), 7.99 (1H, d, J=8.8 Hz), 7.87 (1H, d, J=8.1 Hz), 7.85 (1H, d, J=8.7 Hz), 7.70 (1H, t, J=8.4 Hz), 7.42-7.33 (3H, m), 7.18 (1H, d, J=8.8 Hz), 5.60 (2H, s), 4.11 (1H, m), 3.62-3.54 (4H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	633 (M+1)	

Table 255

Example No.	431	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, s), 8.16 (1H, d, J=8.8 Hz), 7.99 (1H, d, J=8.7 Hz), 7.74-7.60 (4H, m), 7.37 (2H, t, J=8.8 Hz), 7.28 (1H, dd, J=2.2 Hz, 12.2 Hz), 7.14 (1H, dd, J=2.2 Hz, 8.6 Hz), 5.17 (2H, s), 4.10 (1H, m), 3.15 (6H, brs), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.15 (3H, m)
Purity	> 90% (NMR)	
MS	616 (M+1)	

Example No.	432	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.45 (1H, d, J=7.7 Hz), 8.32 (1H, s), 8.19 (1H, d, J=8.8 Hz), 8.02-7.99 (2H, m), 7.70 (1H, t, J=8.6 Hz), 7.60 (2H, dd, J=5.4 Hz, 8.7 Hz), 7.37 (2H, t, J=8.8 Hz), 7.27 (1H, dd, J=2.3 Hz, 12.2 Hz), 7.14 (1H, dd, J=2.2 Hz, 8.7 Hz), 5.16 (2H, s), 4.20-4.00 (2H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m), 1.18 (6H, d, J=6.6 Hz)
Purity	> 90% (NMR)	
MS	630 (M+1)	

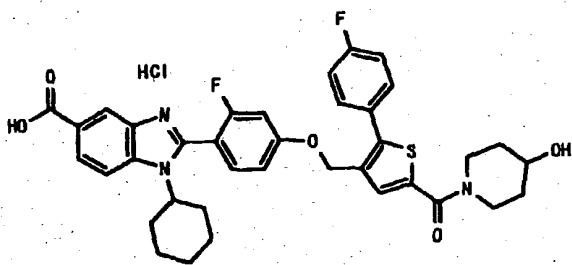
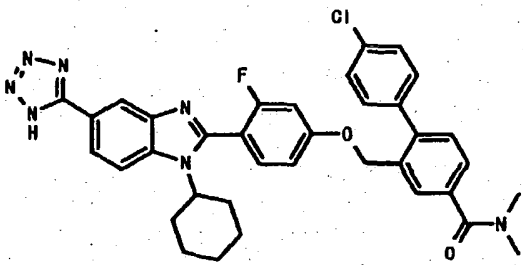
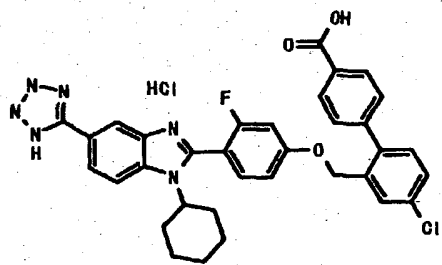
Example No.	433	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.31 (1H, d, J=1.4 Hz), 8.15 (1H, d, J=8.8 Hz), 7.98 (1H, dd, J=1.4 Hz, 8.7 Hz), 7.68-7.60 (4H, m), 7.36 (2H, t, J=8.8 Hz), 7.28 (1H, d, J=2.2 Hz, 12.2 Hz), 7.15 (1H, d, J=2.2 Hz, 8.6 Hz), 5.17 (2H, s), 4.10 (1H, m), 4.05-3.90 (2H, m), 3.85-3.70 (1H, m), 3.55-3.25 (2H, m), 2.40-2.10 (2H, m), 2.00-1.75 (6H, m), 1.70-1.55 (1H, m), 1.50-1.20 (5H, m)
Purity	> 90% (NMR)	
MS	672 (M+1)	

Table 256

Example No.	434	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.45 (1H, d, J=1.5Hz), 8.26 (1H, d, J=8.8Hz), 8.10 (1H, dd, J=8.8, 1.5Hz), 7.72 (1H, d, J=1.5Hz), 7.64 (1H, t, J=8.6Hz), 7.56-7.48 (5H, m), 7.44 (1H, d, J=7.7Hz), 7.18 (1H, dd, J=12.3, 2.4Hz), 7.04 (1H, dd, J=8.6, 2.4Hz), 5.15 (2H, s), 4.08 (1H, brt, J=11.7Hz), 3.02 (3H, s), 2.99 (3H, s), 2.34-2.17 (2H, brm), 1.97-1.81 (4H, brm), 1.70-1.60 (1H, brm), 1.49-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	650 (M+1)	

Example No.	435	1H NMR (δ) ppm
		300Mz, DMSO-d6 8.42 (1H, d, J=1.5Hz), 8.24 (1H, d, J=8.8Hz), 8.08 (1H, dd, J=8.8, 1.5Hz), 8.00 (2H, d, J=8.8Hz), 7.79 (1H, d, J=7.8Hz), 7.62 (1H, t, J=8.4Hz), 7.61-7.55 (3H, m), 7.44 (1H, d, J=8.1Hz), 7.16 (1H, dd, J=12.1, 2.6Hz), 7.02 (1H, dd, J=8.4, 2.6Hz), 5.12 (2H, s), 4.07 (1H, brt, J=12.5Hz), 2.33 (2H, brm), 1.96-1.79 (4H, brm), 1.71-1.61 (1H, brm), 1.49-1.21 (3H, brm)
Purity	> 90% (NMR)	
MS	623 (M+1)	

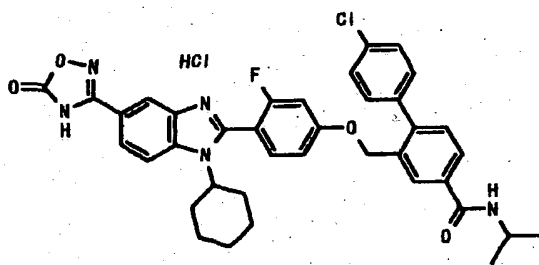
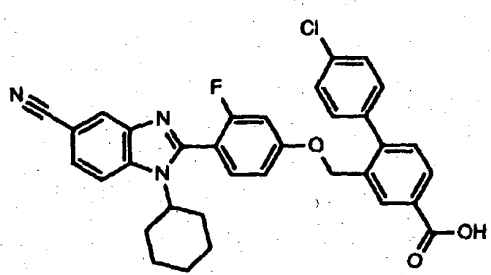
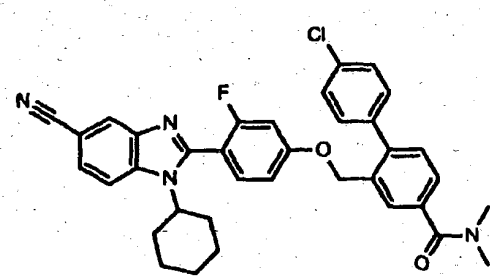
Example No.	436	1H NMR (δ) ppm
		300MHz, DMSO-d6 8.41 (1H, d, J=7.7Hz), 8.30-8.26 (2H, m), 8.18 (1H, d, J=1.4Hz), 7.99 (1H, dd, J=1.7Hz, 8.0Hz), 7.89 (1H, d, J=10.1Hz), 7.67 (1H, t, J=8.8Hz), 7.55-7.45 (5H, m), 7.20 (1H, d, J=12.2Hz), 7.07 (1H, dd, J=2.1Hz, 8.7Hz), 5.14 (2H, s), 4.18-4.11 (2H, m), 2.40-2.10 (2H, m), 2.00-1.75 (4H, m), 1.70-1.55 (1H, m), 1.50-1.20 (3H, m), 1.20 (6H, d, J=6.6Hz)
Purity	> 90% (NMR)	
MS	680 (M+1)	

Table 257

Example No.	437	1H NMR (δ) ppm
		
Purity	> 90% (NMR)	
MS	580 (M+1)	

Example No.	438	1H NMR (δ) ppm
		
Purity	> 90% (NMR)	
MS	607 (M+1)	

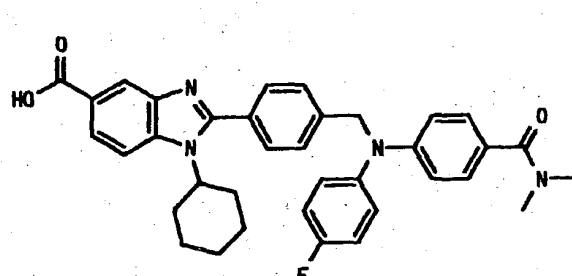
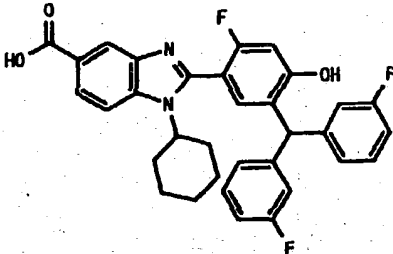
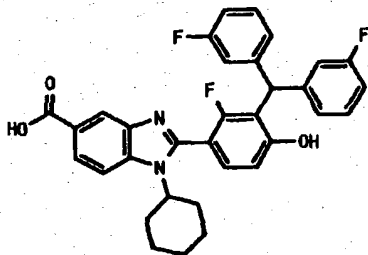
Example No.	439	1H NMR (δ) ppm
		300MHz, CDCl3 8.60 (1H, d, J=1.5Hz), 8.05 (1H, dd, J=1.6Hz, 8.7Hz), 7.70 (1H, d, J=8.7Hz), 7.62 (2H, d, J=8.2Hz), 7.49 (2H, d, J=8.2Hz), 7.31 (2H, d, J=8.8Hz), 7.27-7.23 (2H, m), 7.06 (2H, t, J=8.6Hz), 6.80 (2H, d, J=8.8Hz), 5.05 (2H, s), 4.38 (1H, m), 3.06 (6H, s), 2.45-2.20 (2H, m), 2.10-1.70 (5H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	591 (M+1)	

Table 258

Example No.	440	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.20 (1H, s), 7.86 (2H, m), 7.39 (1H, d, J=7.9Hz), 7.34 (1H, d, J=7.9Hz), 7.07 (2H, dt, J=2.3Hz, 8.6Hz), 6.98-6.88 (5H, m), 6.83 (1H, d, J=8.3Hz), 5.91 (1H, s), 3.96 (1H, m), 2.30-1.95 (2H, m), 1.90-1.50 (4H, m), 1.40-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	557 (M+1)	

Example No.	441	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.24 (1H, d, J=1.4Hz), 8.01 (1H, d, J=8.8Hz), 7.91 (1H, dd, J=1.4Hz, 8.7Hz), 7.47 (1H, t, J=8.4Hz), 7.43-7.35 (2H, m), 7.15-7.01 (5H, m), 6.92 (2H, d, J=10.4Hz), 6.11 (1H, s), 3.90 (1H, m), 2.30-1.95 (2H, m), 1.90-1.50 (4H, m), 1.40-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	557 (M+1)	

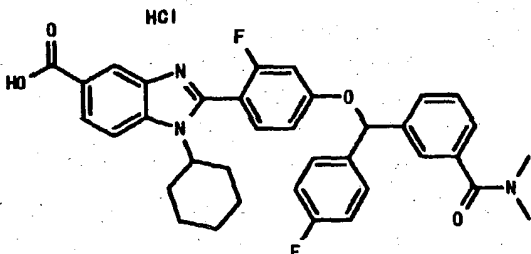
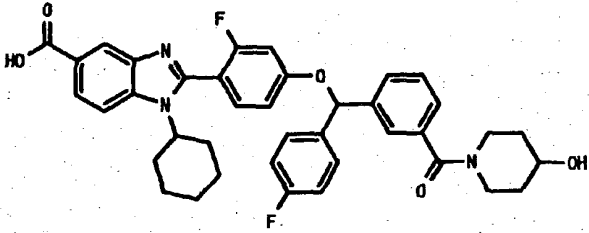
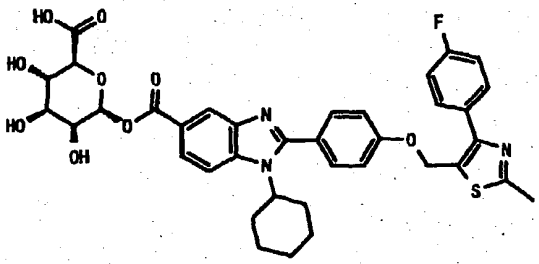
Example No.	442	1H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.26 (1H, d, J=1.5Hz), 8.11 (1H, d, J=8.9Hz), 7.96 (1H, dd, J=8.9, 1.5Hz), 7.65-7.57 (5H, m), 7.47 (1H, t, J=7.7Hz), 7.35 (1H, d, J=7.6Hz), 7.30-7.22 (3H, m), 7.16 (1H, dd, J=8.7, 2.3Hz), 6.88 (1H, s), 4.04 (1H, brt, J=11.3Hz), 2.98 (3H, s), 2.84 (3H, s), 2.30-2.10 (2H, brm), 1.94-1.75 (4H, brm), 1.68-1.57 (1H, brm), 1.45-1.14 (3H, brm)
Purity	> 90% (NMR)	
MS	610 (M+1)	

Table 259 -

Example No.	443	1H NMR (δ) ppm
		300Mz, DMSO-d <sub>6</sub> 8.23 (1H, s), 7.98 and 7.89 (2H, ABq, J=8.8Hz), 7.62-7.06 (11H, m), 6.86 (1H, s), 4.12-3.77 (2H, brm), 3.72 (1H, brs), 3.69 (1H, brs), 3.18 (1H, brs), 3.05 (1H, brs), 2.31-2.08 (2H, brm), 1.90-1.54 (7H, brm), 1.48-1.13 (5H, brm)
Purity	> 90% (NMR)	
MS	666 (M+1)	

Example No.	444	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.36 (1H, s), 8.00 (1H, d, J=8.7Hz), 7.90 (1H, d, J=9.3Hz), 7.80-7.70 (2H, m), 7.63 (2H, d, J=8.4Hz), 7.32 (2H, t, J=8.7Hz), 7.22 (2H, d, J=8.4Hz), 5.62 (1H, d, J=7.5Hz), 5.57 (1H, brd, J=4.8Hz), 5.41 (2H, s), 5.31 (1H, m), 4.29 (1H, m), 3.84 (1H, d, J=9.0Hz), 3.50-3.20 (3H, m), 2.71 (3H, s), 2.40-2.20 (2H, m), 1.75-1.60 (1H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	718 (M+1)	

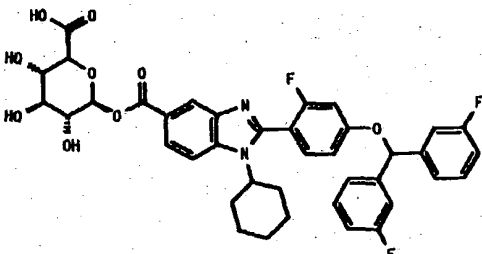
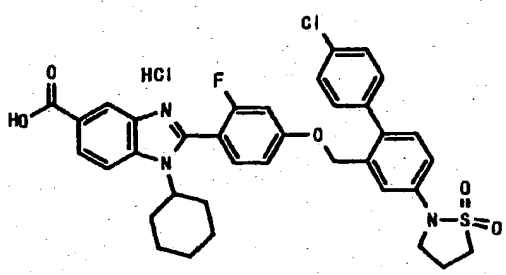
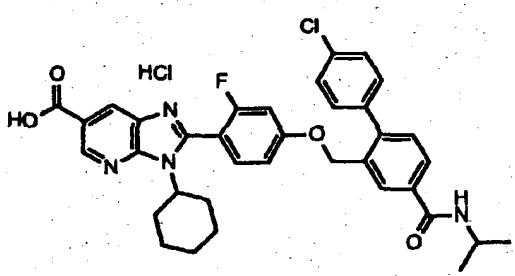
Example No.	445	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.36 (1H, s), 8.00 (1H, d, J=8.7Hz), 7.92 (1H, d, J=9.3Hz), 7.57 (1H, t, J=8.4Hz), 7.50-7.35 (6H, m), 7.25-7.05 (4H, m), 6.82 (1H, s), 5.62 (1H, d, J=7.2Hz), 5.56 (1H, m), 5.28 (1H, brs), 3.95 (1H, m), 3.82 (1H, d, J=8.7Hz), 3.50-3.20 (3H, m), 2.30-2.05 (2H, m), 1.90-1.55 (5H, m), 1.40-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	733 (M+1)	

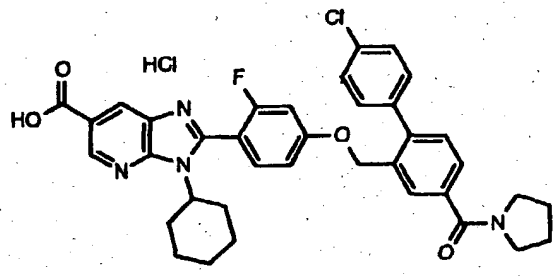
Table 260

Example No.	446	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.29 (1H, s), 8.13 (1H, d, J=9.0Hz), 7.97 (1H, d, J=9.0Hz), 7.63 (1H, t, J=8.6Hz), 7.51-7.32 (7H, m), 7.15 (1H, d, J=12.0Hz), 7.03 (1H, d, J=9.0Hz), 5.10 (2H, s), 4.09 (1H, m), 3.82 (2H, t, J=6.3Hz), 3.56 (2H, t, J=7.4Hz), 2.45 (2H, m), 2.40-2.10 (2H, m), 2.00-1.55 (5H, m), 1.50-1.20 (3H, m)
Purity	> 90% (NMR)	
MS	674 (M+1)	

Example No.	702	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.97 (1H, d, J=1.8Hz), 8.52 (1H, d, J=2.4Hz), 8.36 (1H, d, J=7.8Hz), 8.16 (1H, s), 7.96 (1H, d, J=8.1Hz), 7.55-7.40 (5H, m), 7.14 (1H, d, J=12.6Hz), 7.01 (1H, dd, J=8.4Hz, 1.8Hz), 5.11 (2H, s), 4.20-3.95 (2H, m), 2.65-2.45 (2H, m), 1.95-1.80 (5H, m), 1.20-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	641 (M+1)	

Example No.	703	1H NMR (δ) ppm
		300MHz, DMSO-d <sub>6</sub> 8.97 (1H, d, J=1.8Hz), 8.52 (1H, d, J=1.8Hz), 7.82 (1H, s), 7.70-7.35 (7H, m), 7.13 (1H, d, J=12.3Hz), 7.00 (1H, d, J=11.1Hz), 5.14 (2H, s), 3.60-3.35 (4H, m), 2.65-2.40 (2H, m), 2.00-2.55 (9H, m), 1.40-1.10 (3H, m)
Purity	> 90% (NMR)	
MS	653 (M+1)	

Formulation Example is given in the following. This example is merely for the purpose of exemplification and does not limit the invention.

#### Formulation Example

5	(a) compound of Example 1	10 g
	(b) lactose	50 g
	(c) corn starch	15 g
	(d) sodium carboxymethylcellulose	44 g
	(e) magnesium stearate	1 g

10 The entire amounts of (a), (b) and (c) and 30 g of (d) are kneaded with water, dried in vacuo and granulated. The obtained granules are mixed with 14 g of (d) and 1 g of (e) and processed into tablets with a tableting machine to give 1000 tablets each containing 10 mg of (a).

15

#### Industrial Applicability

As is evident from the above-mentioned results, the compound of the present invention shows a high inhibitory activity against HCV polymerase.

20 Therefore, the compound of the present invention can provide a pharmaceutical agent effective for the prophylaxis or treatment of hepatitis C, based on the anti-HCV effect afforded by the HCV polymerase inhibitory activity. When used concurrently with a different anti-HCV agent, such as interferon, and/or an  
25 anti-inflammatory agent and the like, it can provide a pharmaceutical agent more effective for the prophylaxis or treatment of hepatitis C. Its high inhibitory activity specific to HCV polymerase suggests the possibility of the compound being a pharmaceutical agent with slight side effects, which can be  
30 used safely for humans.

This application is based on patent application Nos. 369008/1999, 391904/2000 and 193786/2001 filed in Japan, and international application No. PCT/JP00/09181, the contents of  
35 which are hereby incorporated by reference.